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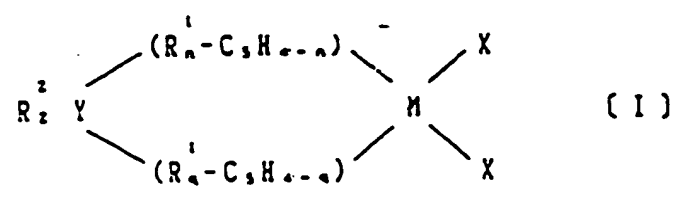
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A transition-metal compound having a bis-substituted, bridged cyclopentadienyl ligand.

Transition-metal compounds have the formula:



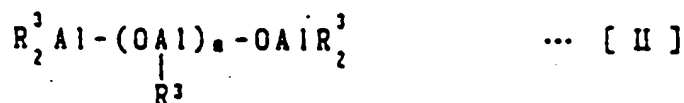
in which M is a zirconium or hafnium atom; Y is a silicon or germanium atom; the groups $\text{R}^n \text{---} \text{C}_5\text{H}_{4-n}$ and $\text{R}^q \text{---} \text{C}_5\text{H}_{4-q}$ each represent an unsubstituted or substituted cyclopentadienyl group, n and q each being 0 or an integer of from 1 to 4 provided that n and q are not simultaneously 0 or 4, the respective substituents R^n and R^q

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EP 0 316 155 A2

being the same or different and each being a hydrogen atom, a silyl group or a hydrocarbon radical; the respective groups R^2 are the same or different and each is hydrogen atom or a hydrocarbon radical; and the respective groups X are the same or different and each is hydrogen or halogen atom or a hydrocarbon radical.

Catalysts for the stereoregulator polymerization of olefins comprise a compound of formula (I) together with an aluminoxane of the formulae:



or



in which m is an integer of from 4 to 30 and R^3 is a hydrocarbyl radical.

A transition-metal compound having a bis-substituted bridged cyclopentadienyl ligand

This invention relates to transition-metal compounds having a bis-substituted-cyclopentadienyl ligand of bridged structure. The compounds of the invention are useful as catalyst components for stereo-regular olefin polymerization.

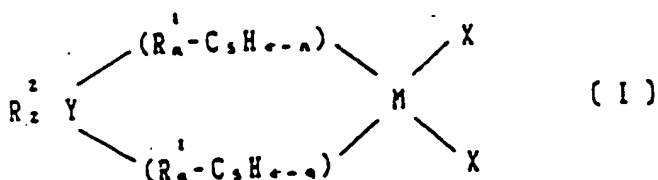
As a homogeneous catalyst for olefin polymerization the so-called "Kaminsky catalyst" (metallocene compound:aluminoxane) is well known. The catalyst has a very high polymerization activity, and in the case of propylene polymerization, it has also been known that it is possible to produce both atactic and isotactic polypropylene using the catalyst [Makromol. Chem. Rapid Commun. 4,417-421 (1983); Japanese laid open patent applications Nos. Sho 58-19,309, Sho 60-130,604 and Sho 61-211,307; Angew. Chem. Soc. 1984, 106, 6355-6364; Japanese laid-open patent application No. Sho 61-264,010 J. Am. Chem. Soc., 1987, 109, 6544-6545; and Japanese laid open patent application No. Sho. 63-66206.

As to metallocene compounds, transition-metal compounds having a bis-unsubstituted-cyclopentadienyl ligand crosslinked with silicon, phosphine, amine or carbon are known (Japanese laid-open patent application No. Sho 62-296,008) and as to zirconium or hafnium compounds, dimethylsilylbis (cyclopentadienyl)-zirconium or hafnium dichloride has been synthesized (Z. Naturforsch. 38b, 321-325 (1983) and Inorg. Chem. 1985, 24, 2539).

Further, as examples of transition-metal compounds having a ligand using substituted-cyclopentadienyl rings crosslinked with silicon, dimethylsilylbis (methylcyclopentadienyl)titanium dichloride (Monatsh. Chem. 112,887-897 (1981)), dimethylsilylbis-(t-butylcyclopentadienyl)scandium hydride (J. Am. Chem. Soc. 1988, 110,976-978), $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5) \text{MCG}[\text{SiMe}_3]_2$ ($\text{M} = \text{Nd, Sm, Lu}$) type compounds (J. Am. Chem. Soc. 1985, 107, 8103-8110), etc. have been described.

With regard to zirconium or titanium compounds, an example of those having a dimethylsilylbis-(tetramethylcyclopentadienyl)ligand is known (Chem. Ber. 119, 1750-1754 (1986)), but the preparation of compounds having a ligand using other substituted-cyclopentadienyl rings crosslinked with silicon or germanium has not been reported. Even less, no concrete practical examples wherein olefins are polymerized using such compounds to obtain isotactic polymers have been reported.

According to one embodiment of the invention there are provided transition-metal compounds of the formula:



in which M is a zirconium or hafnium atom; Y is a silicon or germanium atom; the groups $\text{R}^1 - \text{C}_5\text{H}_{4-n}$ and $\text{R}^1 - \text{C}_5\text{H}_{4-q}$ each represent a substituted or unsubstituted cyclopentadienyl group, n and q each being 0 or an integer of from 1 to 4, provided that n and q are not simultaneously 0 or 4, and respectively substituted R¹ may be the same or different and each is a hydrogen atom, a silyl group or a hydrocarbon radical; the groups R² are the same or different and each is hydrogen atom or a hydrocarbon radical; and the groups X may be the same or different and each is hydrogen or halogen atom or a hydrocarbon radical.

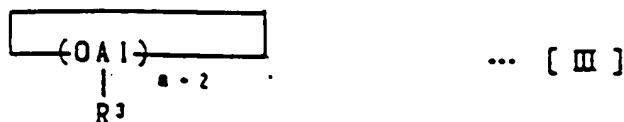
The invention also provides a catalyst for producing a stereoregular olefin polymer, which catalyst comprises as its effective components:

(A) a transition-metal compound of formula (I) as defined above, provided that the position of R¹ on the cyclopentadienyl ring and the nature of R¹ give a configuration wherein any symmetry plane containing M is absent; and

(B) an aluminoxane of the formulae:



or



in which m is an integer of from 4 to 30 and R^3 represents a hydrocarbon radical.

In the accompanying drawings:

Figs. 1, 3, 5, 7, 9, 11, 13 and 15 respectively show the IR spectra of the compounds shown in Table A below;

Figs. 2, 4, 6, 8, 10, 12, 14 and 16 respectively show the NMR spectra of the compounds shown in Table A below

Table A

Zirconium compound	IR spectra	NMR spectra
Me ₂ Si(Me-C ₅ H ₃) ₂ ZrCl ₂	Fig. 1	Fig. 2
Me ₂ Si(C ₅ H ₄)(Me-C ₅ H ₃)ZrCl ₂	Fig. 3	Fig. 4
Me ₂ Si(C ₅ H ₄)(t-Bu-C ₅ H ₃)ZrCl ₂	Fig. 5	Fig. 6
Me ₂ Si(Me-C ₅ H ₃)(t-Bu-C ₅ H ₃)ZrCl ₂	Fig. 7	Fig. 8
Me ₂ Si(t-Bu-C ₅ H ₃) ₂ ZrCl ₂	Fig. 9	Fig. 10
Me ₂ Ge(Me-C ₅ H ₃) ₂ ZrCl ₂	Fig. 11	Fig. 12
(CH ₂) ₄ Si(C ₅ H ₄)(Me-C ₅ H ₃)ZrCl ₂	Fig. 13	Fig. 14
Me ₂ Si(Me-C ₅ H ₃)(Me ₂ -C ₅ H ₂)ZrCl ₂	Fig. 15	Fig. 16

Figs. 17, 19 and 21 respectively show the IR spectra of the compounds shown in Table B below;

Figs. 18, 20 and 22 respectively show the NMR spectra of the compounds shown in Table B below.

Table B

Hafnium compound	IR spectra	NMR spectra
Me ₂ Si(Me-C ₅ H ₃) ₂ HfCl ₂	Fig. 17	Fig. 18
Me ₂ Si(C ₅ H ₄)(Me-C ₅ H ₃)HfCl ₂	Fig. 19	Fig. 20
Me ₂ Si(t-Bu-C ₅ H ₃) ₂ HfCl ₂	Fig. 21	Fig. 22

Fig. 23 is a flowsheet illustrating polymerization using the catalyst of the present invention.

The compounds of formula (I) are zirconium or hafnium compounds having a bidentate ligand of bis-substituted-cyclopentadienyl rings having a bridged structure. The compounds include those of racemic form and meso form depending on the steric configuration of the substituents on the two cyclopentadienyl rings and may have either of these structures. Further, those obtained by mixing the two in optional

proportions are also included.

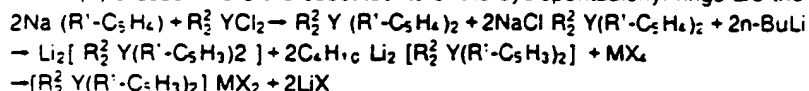
The number of substituents on the respective cyclopentadienyl rings may be from zero to 4, but at least one of the cyclopentadienyl rings is required to have a substituent and also the case where the cyclopentadienyl rings both have 4 substituents is excluded. The respective substituent R' may be the same or different and each is a hydrogen atom, a silyl group or a hydrocarbon radical such as an alkyl, alkenyl, aryl, alkylaryl, arylalkyl groups, etc. each containing up to 20 carbon atoms.

Examples of silyl groups are the trimethylsilyl, triethylsilyl and triphenylsilyl groups. Examples of hydrocarbon radicals are methyl, ethyl, propyl, isopropyl, butyl, sec-butyl, tert-butyl, phenyl, tolyl, biphenyl and naphthyl groups, etc.

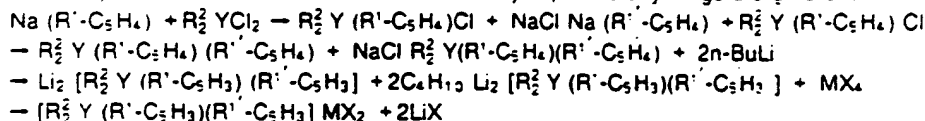
The respective groups R² may be the same or different and each is a hydrogen atom or the above hydrocarbon radical. The respective groups X² may be the same or different and each is a hydrogen atom, a halogen atom such as F, Cl, Br, I, etc. or a hydrocarbon radical as mentioned above.

As to representative preparation routes for the compounds (I) of the invention, the following routes (1) or (2) may be briefly illustrated, for example in the case of n=q-1, but the routes are not limited to these two:

(1) a case where the substituents of the cyclopentadienyl rings are the same:



(2) a case where the substituents of the cyclopentadienyl rings are different:



Preparation of a bidentate ligand of bis-substituted-cyclopentadienyl rings having a bridged structure (e.g. compounds R₂² Y (R'-C₅H₄)₂ or R₂² Y(R'-C₅H₄)(R'-C₅H₄) in the above preparation routes (1) or (2)) has been known (see Monatsh. Chem. 112.887-897 (1981), J. Am. Chem. Soc. 1988, 110.976-978, J. Am. Chem. Soc. 1985, 107, 8103-8110, etc.)

As to such a bidentate ligand, the starting material may be converted with an alkyllithium into a lithium compound, followed by using it for the subsequent reaction. Further, it is also possible to use it in the form of a potassium compound or a sodium compound.

The reaction of MX₄ with a Li compound (e.g. compounds expressed by Li₂ [R₂² Y (R'-C₅H₃)₂] or Li₂ [R₂² Y (R'-C₅H₃) (R'-C₅H₃)] in the above preparation routes (1) and (2)) may be carried out in an ether, preferably tetrahydrofuran (THF) as a reaction solvent. The reaction is preferred to be carried out in a molar ratio of Li compound/MX₄ within a range of 1.0 to 1.5, particularly 1.0 to 1.1. The reaction temperature is preferred to be in a range of -80° to +100° C, particularly 0° to 70° C. The thus formed compound (I) may be purified by recrystallization.

Non-limitative examples of the above-mentioned transition-metal compounds are as follows:

Zirconium compounds

dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride,
diphenylsilylbis(methylcyclopentadienyl)zirconium dichloride,
cyclotrimethylenesilylbis(methylcyclopentadienyl)zirconium dichloride,
cyclotetramethylenesilylbis(methylcyclopentadienyl)zirconium dichloride,
cyclopentamethylenesilylbis(methylcyclopentadienyl)zirconium dichloride,
dimethylsilylbis(methylcyclopentadienyl)zirconium dibromide,
dimethylsilylbis(methylcyclopentadienyl)zirconium methyl chloride,
dimethylsilylbis(methylcyclopentadienyl)zirconium dimethyl,
dimethylsilylbis(methylcyclopentadienyl)zirconium diphenyl
dimethylsilylbis(ethylcyclopentadienyl)zirconium dichloride,
dimethylsilylbis(isopropylcyclopentadienyl)zirconium dichloride,
dimethylsilylbis(tert-butylcyclopentadienyl)zirconium dichloride,
dimethylsilylbis(phenylcyclopentadienyl)zirconium dichloride,
dimethylsilylbis(trimethylsilylcyclopentadienyl)zirconium dichloride,
dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride,
diphenylsilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride.

cycloctetramethylenesilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dimethyl.
 dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium diphenyl.
 dimethylsilyl(cyclopentadienyl) (isopropylcyclopentadienyl)zirconium dichloride.
 5 dimethylsilyl(cyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(cyclopentadienyl) (phenylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(cyclopentadienyl) (trimethylsilylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(methylcyclopentadienyl) (phenylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride.
 10 dimethylgermylbis(methylcyclopentadienyl)zirconium dichloride.
 dimethylgermylbis(t-butylcyclopentadienyl)zirconium dichloride.
 dimethylgermylbis(phenylcyclopentadienyl)zirconium dichloride.
 dimethylgermyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride.
 dimethylgermyl(cyclopentadienyl) (phenylcyclopentadienyl)zirconium dichloride.
 15 dimethylgermyl(cyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride.
 dimethylsilyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)zirconium dichloride.
 dimethylgermyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)zirconium dichloride, etc.

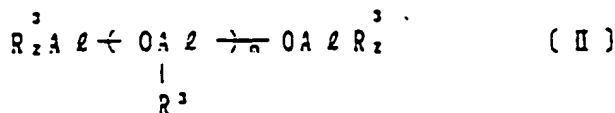
20 Hafnium compounds

dimethylsilylbis(methylcyclopentadienyl)hafnium dichloride.
 diphenylsilylbis(methylcyclopentadienyl)hafnium dichloride.
 cycloctetramethylenesilylbis(methylcyclopentadienyl)hafnium dichloride.
 25 cyclopentamethylenesilylbis(methylcyclopentadienyl)hafnium dichloride.
 cyclopentamethylenesilylbis(methylcyclopentadienyl)hafnium dichloride.
 dimethylsilylbis(methylcyclopentadienyl)hafnium dibromide.
 dimethylsilylbis(methylcyclopentadienyl)hafnium methyl chloride.
 dimethylsilylbis(methylcyclopentadienyl)hafnium dimethyl.
 30 dimethylsilylbis(methylcyclopentadienyl)hafnium diphenyl.
 dimethylsilylbis(ethylcyclopentadienyl)hafnium dichloride.
 dimethylsilylbis(isopropylcyclopentadienyl)hafnium dichloride.
 dimethylsilylbis(t-butylcyclopentadienyl)hafnium dichloride.
 dimethylsilylbis(phenylcyclopentadienyl)hafnium dichloride.
 35 dimethylsilylbis(trimethylsilylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride.
 diphenylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride.
 cycloctetramethylenesilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dimethyl.
 40 dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium diphenyl.
 dimethylsilyl(cyclopentadienyl) (isopropylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(cyclopentadienyl) (t-butylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(cyclopentadienyl) (phenylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(cyclopentadienyl) (trimethylsilylcyclopentadienyl)hafnium dichloride.
 45 dimethylsilyl(methylcyclopentadienyl) (phenylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)hafnium dichloride.
 dimethylgermylbis(methylcyclopentadienyl)hafnium dichloride.
 dimethylgermylbis(t-butylcyclopentadienyl)hafnium dichloride.
 dimethylgermylbis(phenylcyclopentadienyl)hafnium dichloride.
 50 dimethylgermyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride.
 dimethylgermyl(cyclopentadienyl) (phenylcyclopentadienyl)hafnium dichloride.
 dimethylgermyl(cyclopentadienyl) (t-butylcyclopentadienyl)hafnium dichloride.
 dimethylsilyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)hafnium dichloride.
 dimethylgermyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)hafnium dichloride, etc.

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The zirconium or hafnium compounds [I] of the present invention exhibits a superior olefin polymeriza-

tion activity in the presence of an aluminoxane and is useful compounds as a polymerization catalyst component. The aluminoxane as one of the catalyst components used in the polymerization is an organic aluminum compound expressed by the formula [II] or the formula [III]



wherein R^2 represents a hydrocarbon radical such as methyl group, ethyl group, propyl group, butyl group, etc., preferably methyl group or ethyl group and m represents an integer of 4 to 30, preferably 6 or more, more preferably 10 or more. Preparation of such compounds are known. For example, they may be prepared by adding a trialkylaluminum to a hydrocarbon suspension of a compound containing adsorbed water or a salt containing water of crystallization such as copper sulfate hydrate, aluminum sulfate hydrate, etc., followed by reacting the mixture.

Olefins used for polymerization wherein the catalyst of the present invention is used, are α -olefins such as propylene, 1-butene, 4-methyl-1-pentene, 1-hexene, 1-octene, etc. and mixtures of two or more kinds thereof may also be used for polymerization. Further, it is also possible to copolymerize the above-mentioned α -olefins with ethylene. Still further, the catalyst is effective for copolymerization of α -olefins with conjugated or non-conjugated dienes such as butadiene, 1,4-hexadiene, 1,4-pentadiene, 1,7-octadiene, etc., styrene or cyclic olefins such as cyclopropene, cyclobutene, norbornene, dicyclopentadiene, etc.

As to the polymerization process employed in the present invention, either of liquid phase polymerization or gas phase polymerization is possible. As the polymerization solvent for liquid phase polymerization, hydrocarbons capable of dissolving both the components of the zirconium or hafnium compound and the aluminoxane are used, and concretely, aromatic hydrocarbons such as benzene, toluene, o-xylene, m-xylene, p-xylene, ethylbenzene, butylbenzene, mesitylene, naphthalene, etc., preferably toluene and xylene are used. Further, even in the case of solvents incapable of dissolving both the components of the zirconium compound or the hafnium compound and the aluminoxane, if prepolymerization is carried out in aromatic hydrocarbons in advance, polymerization can be carried out.

Examples of such solvents are aliphatic hydrocarbons such as butane, isobutane, pentane, hexane, octane, decane, dodecane, hexadecane, octadecane, etc., alicyclic hydrocarbons such as cyclopentane, methylcyclopentane, cyclohexane, cyclooctane, etc., petroleum fractions such as gasoline, kerosine, gas oil, etc. Among these, aliphatic hydrocarbons are preferred. Further, it is also possible to use liquefied olefins themselves such as liquefied propylene, liquefied butene-1, etc.

As to the catalyst components, the mixture of both the components of the zirconium compound or hafnium compound and the aluminoxane obtained by mixing the components in advance may be fed to the reaction system, or the components may be separately fed to the reaction system. In either of the cases, the concentration and molar ratio of both the components in the reaction system have no particular limitation, but the concentration of the transition metals is preferably in the range of 10^{-2} to 10^{-1} mol/l, and the molar ratio of Al to M is preferably in the range of 100 or more, particularly 1,000 or more.

The olefin pressure in the reaction system has no particular limitation, but it is preferably in the range of the atmospheric pressure to 50 Kg/cm²G. The polymerization temperature also has no particular limitation, but it is usually in the range of -50° to $+230^\circ$ C, preferably -30° to 100° C. Adjustment of the molecular weight at the time of polymerization may be carried out e.g. by choice of temperature or introduction of hydrogen gas.

According to the present invention, a novel zirconium or hafnium compound having a bis-substituted-cyclopentadienyl ligand of a bridged structure is obtained. When the zirconium compound or hafnium compound is used as a catalyst component for olefin polymerization, it is possible to obtain a stereoregular polymer with a good efficiency.

The present invention will be described in more detail by way of Examples.

Example 1

[Dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride]

In this Example, reactions were all carried out in an inert gas atmosphere. Further, the reaction solvent used was dried in advance. In a 200 ml capacity glass reaction vessel, dimethylbis-
 5 (methylcyclopentadienyl)silane (3.5 g, 16 mmol) was dissolved in tetrahydrofuran (50 ml), followed by gradually dropwise adding a hexane solution (23 ml) of n-butyllithium (1.6 M) (M refers to mol l) to the above solution under ice-cooling and agitating the mixture at room temperature for 2 hours to obtain a yellow transparent liquid ($\text{Li}_2[\text{Me}_2\text{Si}(\text{MeC}_5\text{H}_3)_2]$).

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (3.5 g, 15 mmol) was cooled down
 10 to -78°C , followed by adding tetrahydrofuran (200 ml), thereafter gradually dropwise adding the yellow transparent liquid obtained above, at -78°C , agitating the mixture at room temperature for 15 hours, distilling off the solvent from a yellow-orange color solution containing white precipitates, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating two layers, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding
 15 pentane, filtering off the resulting solids to obtain pale green solids (2.87 g), and recrystallizing the solids from hot heptane (400 ml) to obtain yellow-white crystals (2.02 g). The values of the physical properties of this compound are shown below. In addition, IR spectra were measured according to KBr method and ^1H -NMR spectra were measured using CDCl_3 as solvent.

IR spectra: shown in Fig. 1.

20 NMR spectra: shown in Fig. 2.

Elemental analysis value: $(\text{C}_{12}\text{H}_{13}\text{Cl}_2\text{SiZr})$				
Calculated value (%):	C:	44.66,	H:	4.82
Observed value (%):	C:	45.02,	H:	4.91

25 Example 2

[Polymerization]

Into a 1.5t capacity SUS autoclave sufficiently purged with nitrogen gas were successively fed purified
 35 toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride (0.005 mmol); followed by raising the temperature up to 30°C , continuously introducing propylene so as to keep the total pressure at 3 $\text{Kg cm}^{-2}\text{G}$ to carry out polymerization for 2 hours, thereafter decomposing the catalyst component with methanol and drying the resulting polypropylene to obtain isotactic polypropylene (130 g). The catalyst
 40 activity was 142 $\text{Kg g Zr}\cdot\text{hr}$ and its molecular weight was 9,400.

45 Example 3

[Polymerization]

Into a 1.5t capacity SUS autoclave sufficiently purged with nitrogen gas were successively fed purified
 50 toluene (100 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride (0.005 mmol), followed by raising the temperature up to 30°C , then continuously introducing propylene so as to keep the total pressure at 1 $\text{Kg cm}^{-2}\text{G}$, carrying out polymerization for 10 minutes, stopping the feed of propylene, purging propylene, adding purified hexane (1,000 ml), raising the temperature up to 30°C , continuously feeding propylene so
 55 as to keep the total pressure at 3 $\text{Kg cm}^{-2}\text{G}$, carrying out polymerization for 2 hours, thereafter decomposing the catalyst component with methanol, drying the resulting polypropylene to obtain isotactic polypropylene (125 g). The catalyst activity was 137 $\text{Kg g Zr}\cdot\text{hr}$.

Comparative example 1

[Polymerization]

Example 2 was repeated except that dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride was replaced by dimethylsilylbis(cyclopentadienyl)zirconium dichloride to obtain atactic polypropylene (29 g). The catalyst activity was 32 Kg.g Zr^{orr} and its molecular weight was 5,300.

Comparative example 2

[Polymerization]

Example 2 was repeated except that dimethylsilylbis(methylcyclopentadienyl)zirconium dichloride was replaced by bis(methyl-t-butylcyclopentadienyl)zirconium dichloride to obtain atactic polypropylene (2.6 g). The catalyst activity was 2.9 Kg.g Zr^{orr}.

Comparative example 3

[Polymerization]

Into a 1.5t capacity SUS autoclave sufficiently purged with nitrogen gas were successively added purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 909) (6.3 mmol) and bis(methylcyclopentadienyl)zirconium dichloride (0.02 mmol), followed by raising the temperature up to 50 °C, continuously introducing propylene into the mixture so as to keep the total pressure at 8 Kg.cm²G, carrying out polymerization for 4 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain atactic polypropylene (220 g). The catalyst activity was 30 Kg.g Zr^{orr}.

Example 4

[Dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used.

In a 200 ml capacity glass reaction vessel, dimethyl (cyclopentadienyl) (methylcyclopentadienyl)silane (3.4 g, 17 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a solution (24 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 2 hours, distilling off the solvent to obtain white solids (Li₂[Me₂Si(C₅H₄) (Me-C₅H₃)]), and adding tetrahydrofuran (50 ml) thereto to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (4.0 g, 17 mmol) was cooled down to -78 °C, followed by adding tetrahydrofuran (200 ml), then slowly dropwise adding the yellow transparent liquid obtained above at -78 °C, agitating the mixture at room temperature for one hour, heating it under reflux for one hour, allowing it to cool down, distilling off the solvent from a pale yellow solution containing white precipitates, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering off the resulting solids to obtain pale green solids (2.44 g) and recrystallizing them from hot heptane (400 ml) to obtain pale yellow crystals (1.18 g). The values of the physical properties of this compound are shown below. In addition, IR spectra were measured according to KBr method and ¹H NMR spectra were measured using CDCl₃ as solvent.

Elemental analysis value: (C ₁₃ H ₁₆ Cl ₂ SiZr)				
Calculated value (%):	C:	43.08.	H:	4.45
Observed value (%):	C:	42.86.	H:	4.49

IR spectra: shown in Fig. 3.
NMR spectra: shown in Fig. 4

Example 5

[Polymerization]

Into a 1.5l capacity SUS autoclave sufficiently purged with nitrogen gas were successively added purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilyl(cyclopentadienyl)(methylcyclopentadienyl)zirconium dichloride (0.005 mmol), followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G to carry out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (70 g). The catalyst activity was 77 Kg/g Zr·hr. Its molecular weight was 6,600.

Example 6

[Dimethylsilyl(cyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, dimethyl(cyclopentadienyl) (t-butylcyclopentadienyl)silane (5.1 g, 21 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (29 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 2 hours, distilling off the solvent to obtain white solids (Li₂[Me₂Si(C₅H₄) (t-Bu-C₅H₃)])) and adding thereto tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

In a 500 ml glass reaction vessel, zirconium tetrachloride (5.0 g, 21 mmol) was cooled down to -78 °C, followed by adding tetrahydrofuran (200 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78 °C, agitating the mixture at room temperature for 2 hours, heating it under reflux for 2 hours, allowing it to cool down, distilling off the solvent from the resulting pale yellow solution containing white precipitates, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering off the resulting solids to obtain pale green solids and recrystallizing them from hot heptane (400 ml) to obtain pale yellow crystals (0.8 g). The values of the physical properties of the compound are shown below. In addition, its IR spectra were measured according to KBr method and ¹H-NMR spectra were measured using CDCl₃ as solvent.

Elemental analysis value: (C ₁₆ H ₂₂ Cl ₂ SiZr)				
Calculated value (%):	C:	47.50.	H:	5.48
Observed value (%):	C:	47.65.	H:	5.61

IR spectra: shown in Fig. 5.
NMR spectra: shown in Fig. 6.

Example 7

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas were successively added purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride (0.005 mmol), followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (75 g). The catalyst activity was 82 Kg/g Zr·hr. Further, its molecular weight was 4,300.

Example 8

[Dimethylsilyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, dimethyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)silane (3.9 g, 15 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (20 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 2 hours, distilling off the solvent to obtain white solids (Li₂[Me₂Si(Me-C₅H₇) (t-Bu-C₅H₇)] and adding thereto tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (3.5 g, 15 mmol) was cooled down to -78 °C, followed by adding tetrahydrofuran (200 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78 °C, agitating the mixture at room temperature for 18 hours, heating it under reflux for 4 hours, allowing it to cooling down, distilling off the solvent from a pale yellow solution containing white precipitates, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml), separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane and filtering to obtain pale yellow crystals (0.37 g) from the pale yellow filtrate. The values of the physical properties of this compound are shown below. In addition, the IR spectra were measured according to KBr method and the ¹H-NMR spectra were measured using CDCl₃ as solvent.

Elemental analysis values: (C ₁₇ H ₂₄ Cl ₂ SiZr)				
Calculated value (%):	C:	48.78,	H:	5.78
Observed value (%):	C:	48.81,	H:	5.82

IR spectra: shown in Fig. 7.
NMR spectra: shown in Fig. 8.

Example 9

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas were successively added purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilyl(methylcyclopentadienyl) (t-butylcyclopentadienyl)zirconium dichloride (0.005 mmol), followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (59 g). The catalyst activity was 65 Kg/g Zr·hr. Further, its molecular weight was 6,400.

Example 10

[Dimethylsilylbis(t-butylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, dimethylbis(t-butylcyclopentadienyl)silane (4.6 g, 15 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (21 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 4 hours, distilling off the solvent to obtain white solids ($\text{Li}_2[\text{Me}_2\text{Si}(\text{t-Bu-C}_5\text{H}_7)_2]$) and adding tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (3.5 g, 15 mmol) was cooled down to -78°C , followed by adding tetrahydrofuran (250 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78°C , allowing the mixture to stand, returning the temperature to room temperature, heating it under reflux for 84 hours, allowing the resulting material to cool down, distilling off the solvent from a yellow transparent solution, adding methylene chloride (300 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane and filtering to obtain yellow-white crystals (2.35 g) from a yellow filtrate. The values of the physical properties of this compound are shown below. In addition, the IR spectra were measured according to KBr method and the $^1\text{H-NMR}$ spectra were measured using CDCl_3 as solvent.

Elemental analysis values: ($\text{C}_{20}\text{H}_{30}\text{Cl}_2\text{SiZr}$)				
Calculated value (%):	C:	52.15,	H:	6.56
Observed value (%):	C:	52.01,	H:	6.58

IR spectra: shown in Fig. 9.

NMR spectra: shown in Fig. 10.

Example 11

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas were successively added purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(t-butylcyclopentadienyl)zirconium dichloride (0.005 mmol), followed by raising the temperature up to 30°C , continuously introducing propylene so as to keep the total pressure at $3\text{ Kg}\cdot\text{cm}^{-2}\text{G}$, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (3.1 g). The catalyst activity was $3.4\text{ Kg/g Zr}\cdot\text{hr}$. Its molecular weight was 9,600.

Example 12

[Dimethylgermylbis(methylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, dimethylbis(methylcyclopentadienyl)germane (1.2 g, 8 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (11 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 2 hours, distilling off the solvent to obtain white solids ($\text{Li}_2[\text{Me}_2\text{Ge}(\text{Me-C}_5\text{H}_7)_2]$), adding thereto tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (1.8 g, 8 mmol) was cooled down to -78°C , followed by adding tetrahydrofuran (150 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78°C , agitating the mixture at room temperature for 14 hours, heating it under reflux for 2 hours, allowing the resulting material to cool down, distilling off the solvent from a yellow transparent solution, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering and concentrating to obtain yellow-white crystals (0.31 g) from

a pale yellow filtrate. The values of the physical properties of this compound are shown below. In addition, the IR spectra were measured according to KBr method and $^1\text{H-NMR}$ spectra were measured using CDCl_3 as solvent.

Elemental analysis values: $(\text{C}_{14}\text{H}_{12}\text{Cl}_2\text{GeZr})$				
Calculated value (%):	C:	39.94.	H:	4.31
Observed value (%):	C:	40.38.	H:	4.38

IR spectra: shown in Fig. 11.
NMR spectra: shown in Fig. 12.

Example 13

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylgermylbis(methylcyclopentadienyl)zirconium dichloride (0.005 mmol) were successively added, followed by raising the temperature up to 30°C , continuously introducing propylene so as to keep the total pressure at $3\text{ Kg}\cdot\text{cm}^2\text{G}$, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (2.5 g). The catalyst activity was 27 Kg/g Zr·hr. Its molecular weight was 7,600.

Example 14

[Cyclotetramethylenesilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, cyclotetramethylene(cyclopentadienyl) (methylcyclopentadienyl)silane (3.4 g, 15 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (22 ml) of *n*-butyllithium (1.6 M), agitating the mixture at room temperature for 3 hours, distilling off the solvent to obtain white solids ($\text{Li}_2[(\text{CH}_2)_4\text{Si}(\text{C}_5\text{H}_4)(\text{MeC}_5\text{H}_3)]$), and adding thereto tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, zirconium tetrachloride (3.5 g, 15 mmol) was cooled down to -78°C , adding tetrahydrofuran (250 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78°C , agitating the mixture at room temperature for 48 hours, distilling off the solvent from a yellow solution containing white precipitates, adding methylene chloride (200 ml), filtering off the resulting white solids, concentrating the resulting yellow filtrate, adding pentane and cooling the mixture down to -30°C to obtain yellow-white crystals (0.39 g). The values of the physical properties of this compound are shown below. In addition, its IR spectra were measured according to KBr method and its $^1\text{H-NMR}$ spectra were measured using CDCl_3 as solvent.

Elemental analysis values: $(\text{C}_{15}\text{H}_{12}\text{Cl}_2\text{SiZr})$				
Calculated value (%):	C:	46.37.	H:	4.67
Observed value (%):	C:	47.03.	H:	4.82

IR spectra: shown in Fig. 13.
NMR spectra: shown in Fig. 14.

Example 15

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and
 5 cyclotetramethylenesilyl(cyclopentadienyl) (methylcyclopentadienyl)zirconium dichloride (0.005 mmol) were successively added, followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (3.6 g). The catalyst activity was 3.9 Kg/g Zr·hr. Its molecular weight was 4,200.

10

Example 16

15 [Dimethylsilyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)zirconium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml glass reaction vessel, dimethyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)silane (3.5 g, 15 mmol) was dissolved in pentane (50 ml), followed by slowly
 20 dropwise adding a hexane solution (22 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 5 hours, distilling off the solvent to obtain white solids (Li₂[Me₂Si(Me-C₅H₃) (Me₂-C₅H₂)]), and adding thereto tetrahydrofuran (50 ml) to obtain a yellow-brown transparent liquid.

In a 500 ml glass reaction vessel, zirconium tetrachloride (3.5 g, 15 mmol) was cooled down to -78 °C, followed by adding tetrahydrofuran (250 ml), slowly dropwise adding the yellow-brown transparent liquid
 25 obtained above at -78 °C, agitating the mixture at room temperature for 18 hours, heating it under reflux for 6 hours, distilling off the solvent from the resulting yellow solution, adding methylene chloride (300 ml), filtering off the resulting white solids, concentrating the resulting yellow filtrate, adding pentane, filtering the resulting solids and recrystallizing the solids from hot heptane (200 ml) to obtain yellow-white crystals (1.08 g). The values of the physical properties of this compound are shown below. In addition, its IR spectra were
 30 measured according to KBr method and its ¹H-NMR spectra were measured using CDCl₃ as solvent.

35

Elemental analysis values: (C ₁₅ H ₂₀ Cl ₂ SiZr)				
Calculated value (%):	C:	46.13,	H:	5.16
Observed value (%):	C:	46.22,	H:	5.16

IR spectra: shown in Fig. 15.

NMR spectra: shown in Fig. 16.

40

Example 17

45 [Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and
 50 dimethylsilyl(methylcyclopentadienyl) (dimethylcyclopentadienyl)zirconium dichloride (0.005 mmol) were successively added, followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol and drying the resulting polypropylene to obtain isotactic polypropylene (45 g). The catalyst activity was 49 Kg/g Zr·hr. Its molecular weight was 5,600.

55

Example 18

[Dimethylsilylbis(methylcyclopentadienyl)hafnium dichloride]

All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml glass reaction vessel, dimethylbis(methylcyclopentadienyl)silane (3.3 g, 15 mmol) was dissolved in tetrahydrofuran (50 ml), followed by slowly dropwise adding a hexane solution (21 ml) of n-butyllithium (1.6 M) under ice cooling and agitating the mixture at room temperature for 2 hours to obtain a yellow transparent liquid ($\text{Li}_2[\text{Me}_2\text{Si}(\text{MeC}_5\text{H}_3)_2]$).

In a 500 ml capacity glass reaction vessel, hafnium tetrachloride (4.8 g, 15 mmol) was cooled down to -78°C , followed by adding tetrahydrofuran (200 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78°C , agitating the mixture at room temperature for 17.5 hours, heating it under reflux for one hour, allowing the resulting material to cool down, distilling off the solvent from a yellow-orange color solution containing white precipitates, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml), separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering off insoluble solids, cooling a yellow filtrate to obtain yellow-white crystals (2.02 g). The values of the physical properties of this compound are shown below. In addition, its IR spectra were measured according to KBr method and its ^1H -NMR spectra were measured using CDCl_3 as solvent.

IR spectra: shown in Fig. 17.
NMR spectra: shown in Fig. 18.

20 Example 19

[Polymerization]

25 Into a 1.5 l capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(methylcyclopentadienyl)hafnium dichloride (0.005 mmol) were successively added, followed by raising the temperature up to 30°C , continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (16 g). The catalyst activity was 8.9 Kg/g Hf·hr, and its molecular weight was 66,800.

35 Example 20

[Dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride]

40 All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml glass reaction vessel, dimethyl(cyclopentadienyl) (methylcyclopentadienyl)silane (3.0 g, 15 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution (20 ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 2 hours and distilling off the solvent to obtain white solids ($\text{Li}_2[\text{Me}_2\text{Si}(\text{C}_5\text{H}_4)(\text{Me-C}_5\text{H}_3)]$), and adding thereto tetrahydrofuran (50 ml) to obtain a yellow transparent liquid.

45 In a 500 ml capacity glass reaction vessel, hafnium tetrachloride (4.8 g, 15 mmol) was cooled down to -78°C , followed by adding tetrahydrofuran (200 ml), slowly dropwise adding the yellow transparent liquid obtained above at -78°C , agitating the mixture at room temperature for 14 hours, distilling off the solvent from a yellow solution containing white solids, adding methylene chloride (200 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering off the resulting solids to obtain pale green solids (2.22 g) and recrystallizing the crystals from n-heptane (200 ml) to obtain pale yellow-white crystals (0.94 g). The values of the physical properties of this compound are shown below. In addition, its IR spectra were measured according to KBr method and its ^1H -NMR spectra were measured using CDCl_3 as solvent.

55 IR spectra: shown in Fig. 19.
NMR spectra: shown in Fig. 29

Example 21[Polymerization]

5 Into a 1.5 l capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilyl(cyclopentadienyl) (methylcyclopentadienyl)hafnium dichloride (0.005 mmol) were succe-
 10 ssively added, followed by raising the temperature up to 30 °C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol and drying the resulting polypropylene to obtain isotactic polypropylene (0.6 g). The catalyst activity was 0.34 Kg/g Hf·hr, and its molecular weight was 52,800.

15 Example 22[Dimethylsilylbis(t-butylcyclopentadienyl)hafnium dichloride]

20 All the reactions were carried out in an inert gas atmosphere. Further, the reaction solvent was dried in advance and used. In a 200 ml capacity glass reaction vessel, dimethylbis(t-butylcyclopentadienyl)silane (4.8 g, 16 mmol) was dissolved in pentane (50 ml), followed by slowly dropwise adding a hexane solution- (22ml) of n-butyllithium (1.6 M) under ice cooling, agitating the mixture at room temperature for 4 hours, distilling off the solvent to obtain white solids (Li₂[Me₂Si(t-Bu-C₅H₃)₂], and adding thereto tetrahydrofuran
 25 (60 ml) to obtain a yellow transparent liquid.

In a 500 ml capacity glass reaction vessel, hafnium tetrachloride (5.1 g, 16 mmol) was cooled down to -78 °C, followed by adding tetrahydrofuran (250 ml), slowly dropwise adding the yellow transparent liquid at -78 °C, allowing the mixture to stand to return the temperature to room temperature, heating it under reflux for 84 hours, allowing the resulting material to cool down, distilling off the solvent from a yellow-orange
 30 color transparent solution, adding methylene chloride (300 ml) and then dilute hydrochloric acid (100 ml) under ice cooling, separating the liquids, drying the methylene chloride layer over anhydrous sodium sulfate, distilling off methylene chloride, adding pentane, filtering, concentrating the resulting yellow filtrate and cooling to obtain pale yellow crystals (1.41 g). The values of the physical properties of this compound are shown below. In addition, its IR spectra were measured according to KBr method and its ¹H-NMR
 35 spectra were measured using CDCl₃ as solvent.

Elemental analysis values: (C ₂₂ H ₂₀ C (t ₂ SiHf)				
Calculated value (%):	C:	43.84.	H:	5.52
Observed value (%):	C:	44.05.	H:	5.62

IR spectra: shown in Fig. 21.

NMR spectra: shown in Fig. 22.

45 Example 23[Polymerization]

50 Into a 1.5 l capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(t-butylcyclopentadienyl)hafnium dichloride (0.005 mmol) were successively added, followed
 55 by raising the temperature up to 30 °C, continuously introducing propylene-so as to keep the total pressure at 3 Kg cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain isotactic polypropylene (0.3 g). The catalyst activity was 0.17 Kg/g Hf·hr. Further, its molecular weight was 17,200.

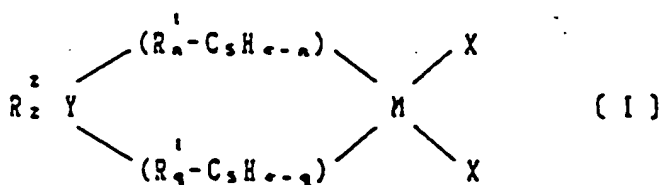
Comparative example 4

[Polymerization]

Into a 1.5 t capacity SUS autoclave sufficiently purged with nitrogen gas, purified toluene (500 ml), methylaluminoxane made by Toyo Stauffer Chemical Co., Ltd. (molecular weight: 770) (4.0 mmol) and dimethylsilylbis(cyclopentadienyl)hafnium dichloride (0.005 mmol) were successively added, followed by raising the reaction up to 30°C, continuously introducing propylene so as to keep the total pressure at 3 Kg/cm²G, carrying out polymerization for 2 hours, decomposing the catalyst component with methanol after the reaction and drying the resulting polypropylene to obtain atactic polypropylene (3.5 g). The catalyst activity was 2.0 Kg/g Hf·hr.

Claims

1. Transition-metal compounds of the formula:

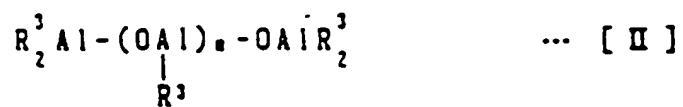


in which M is a zirconium or hafnium atom; Y is a silicon or germanium atom; the groups R'_n-C₅H_{4-n} and R'_q-C₅H_{4-q} each represent an unsubstituted or substituted cyclopentadienyl group, n and q each being 0 or an integer of from 1 to 4 provided that n and q are not simultaneously 0 or 4, the respective substituents R' being the same or different and each being a hydrogen atom, a silyl group or a hydrocarbon radical; the respective groups R² are the same or different and each is hydrogen atom or a hydrocarbon radical; and the respective groups X are the same or different and each is hydrogen or halogen atom or a hydrocarbon radical.

2. A catalyst for producing a stereoregular olefin polymer, which catalyst comprises as its effective components.

(A) a transition-metal compound of formula (I) as defined in claim 1 provided that the site of R' on the cyclopentadienyl ring and the nature of R' give a configuration wherein any symmetry plane containing M is absent; and

(B) an aluminoxane of the formulae:



or

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in which m is an integer of from 4 to 30 and R^3 is a hydrocarbon radical.

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FIG. 1

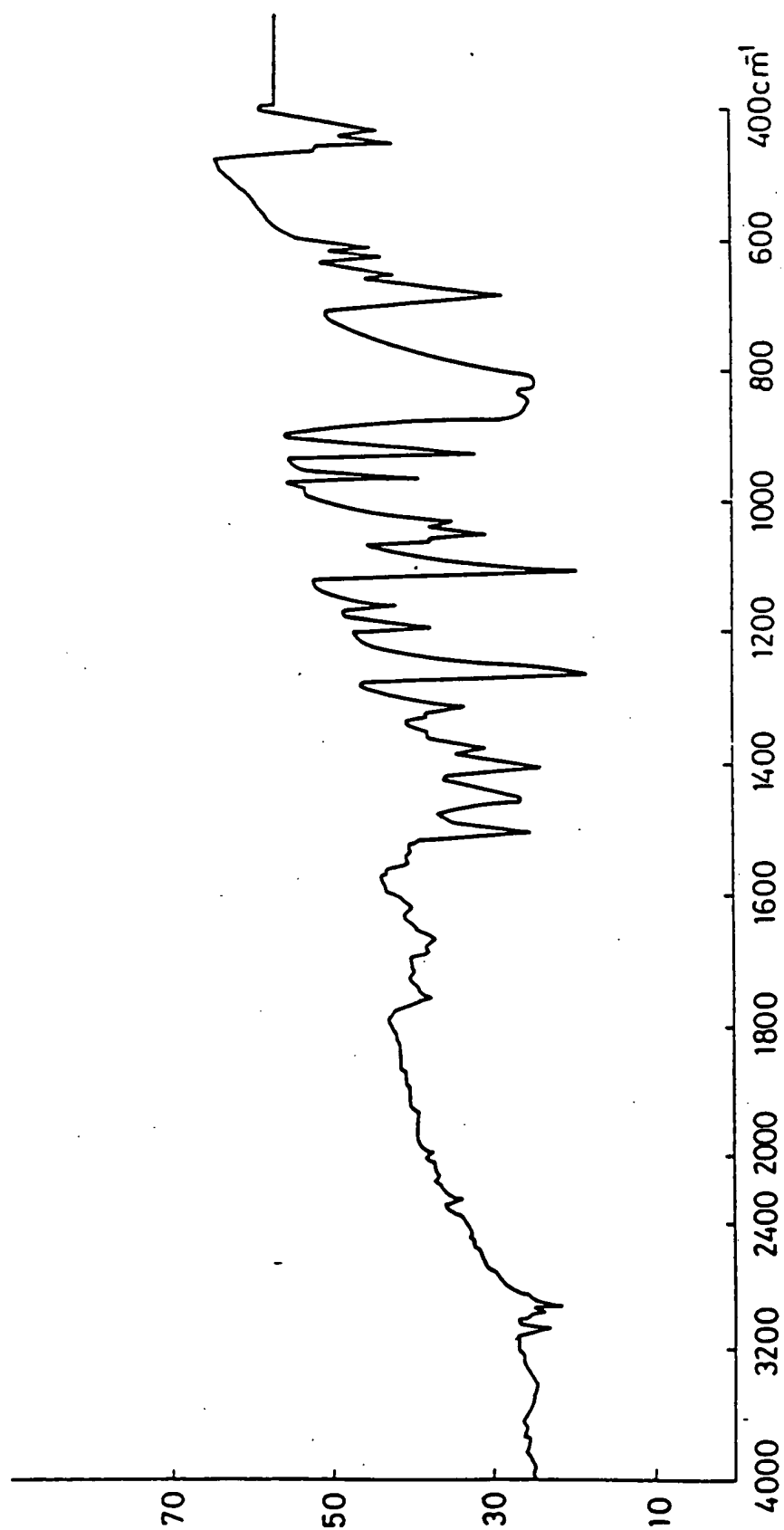


FIG. 2

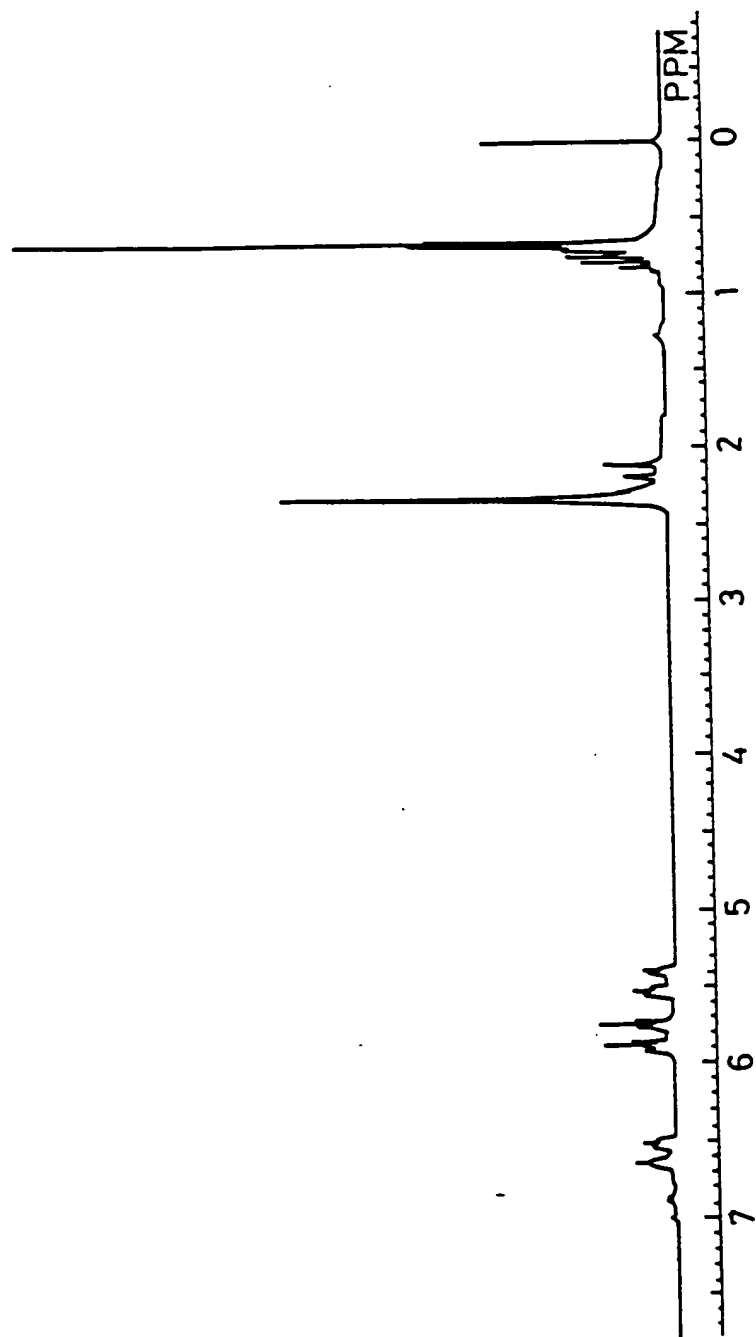


FIG. 3

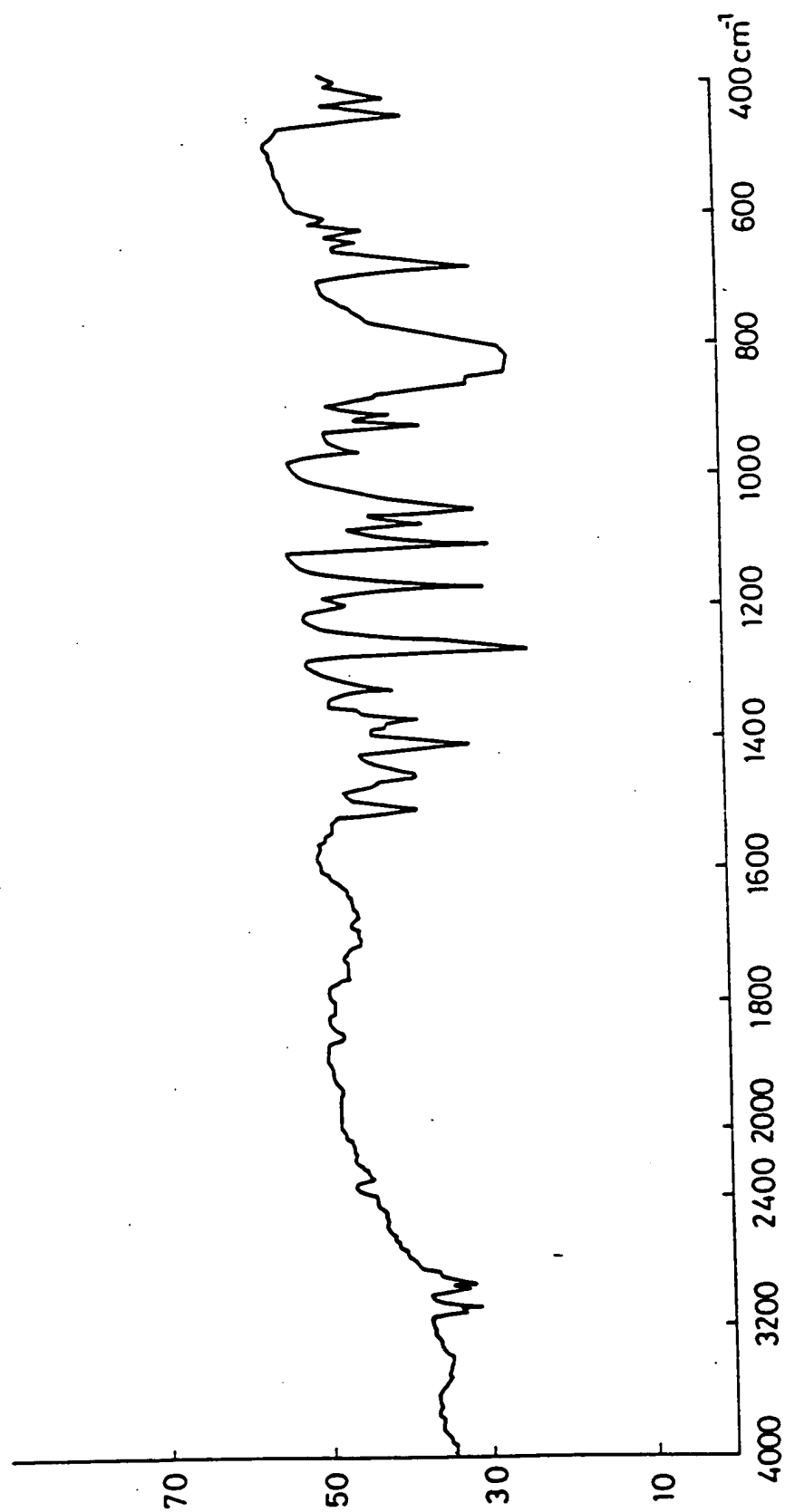


FIG. 4

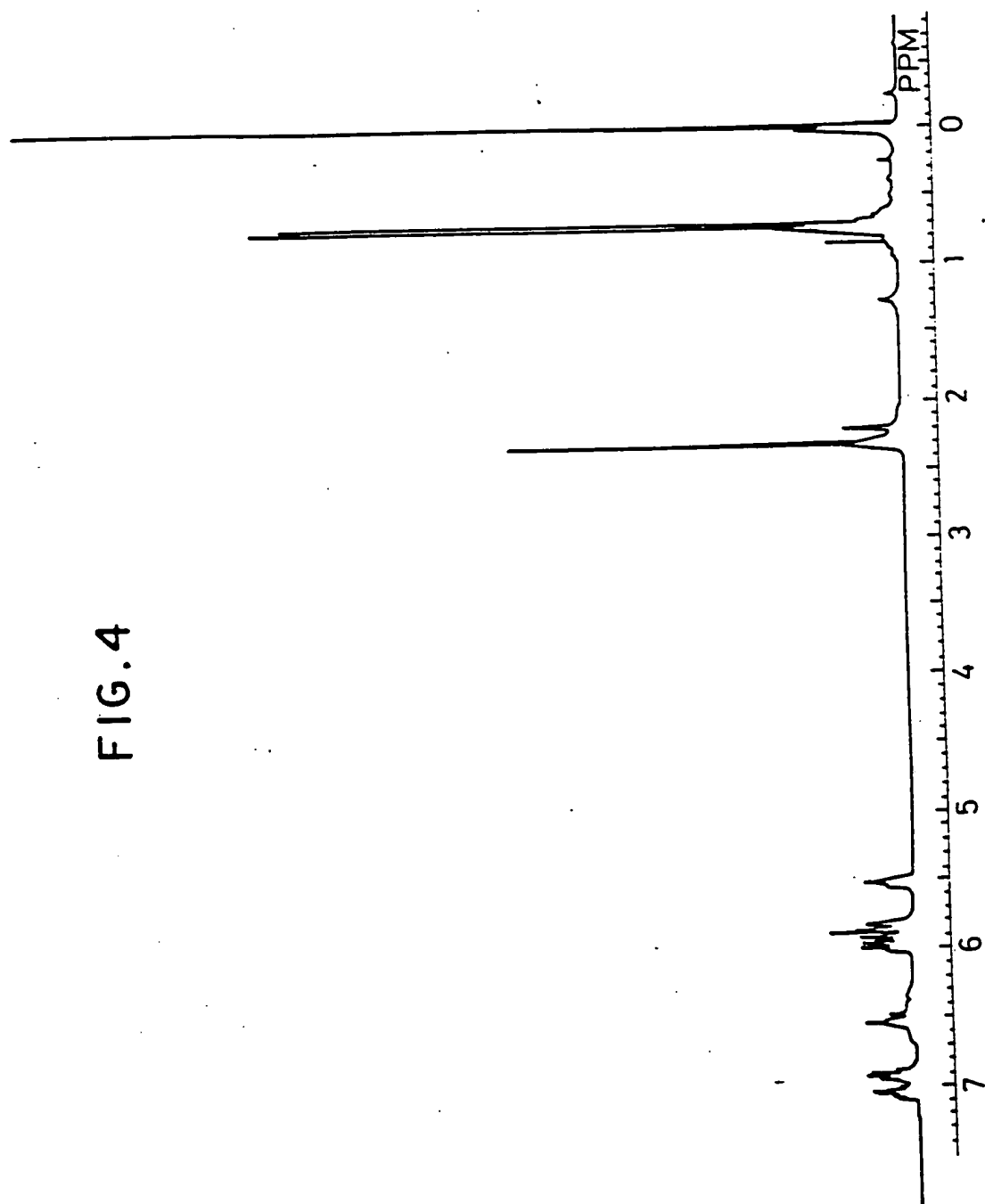


FIG. 5

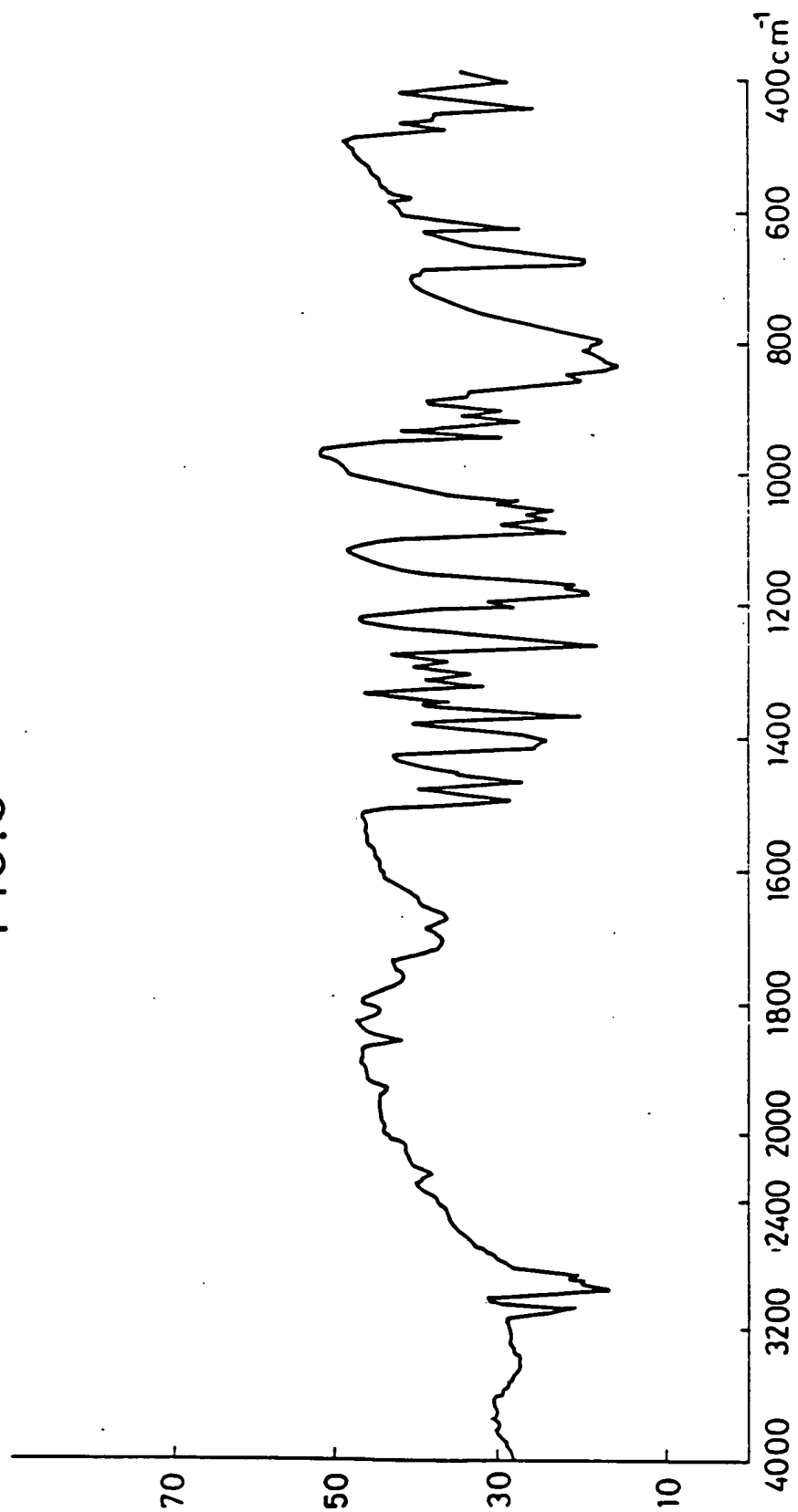


FIG. 6

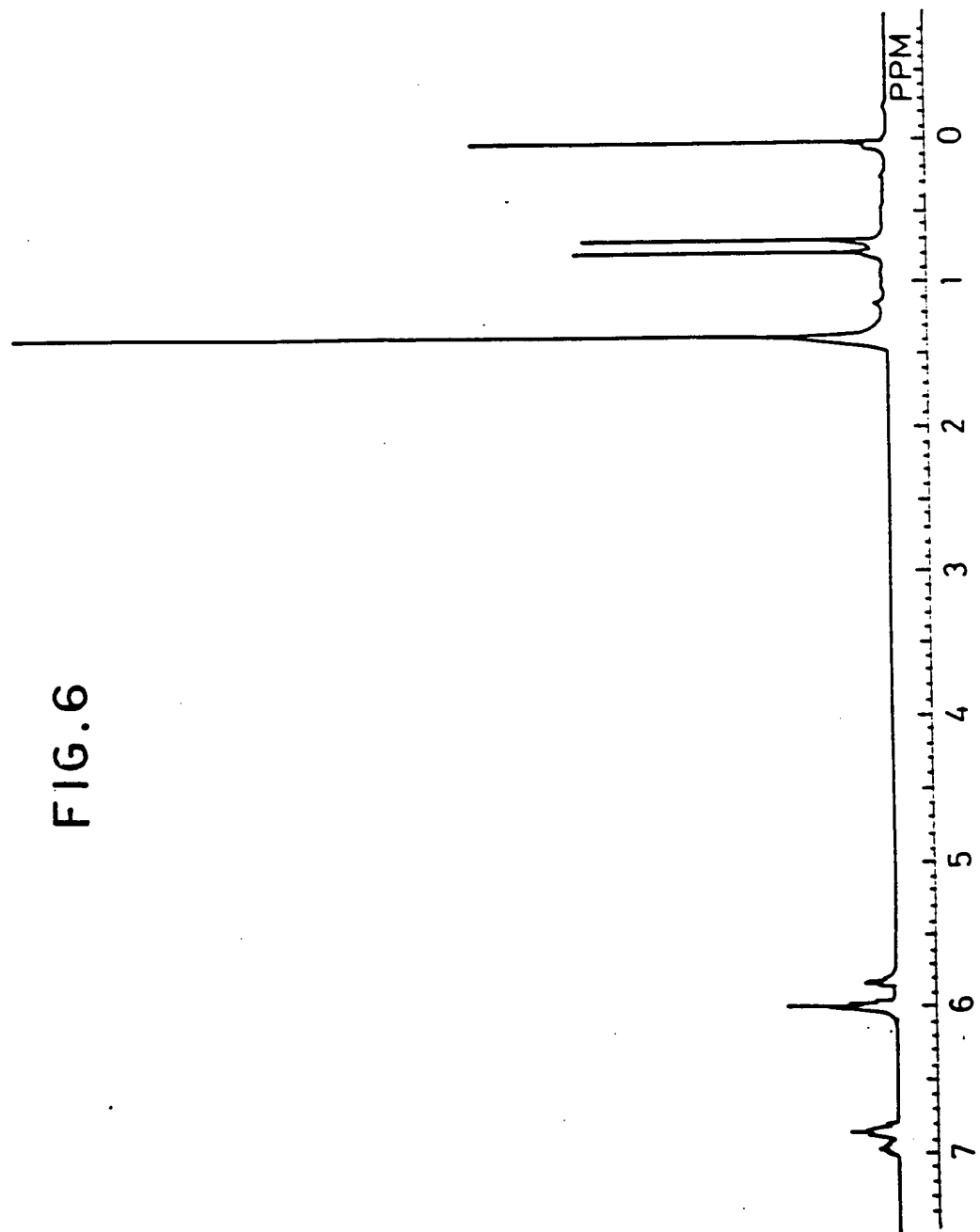


FIG. 7

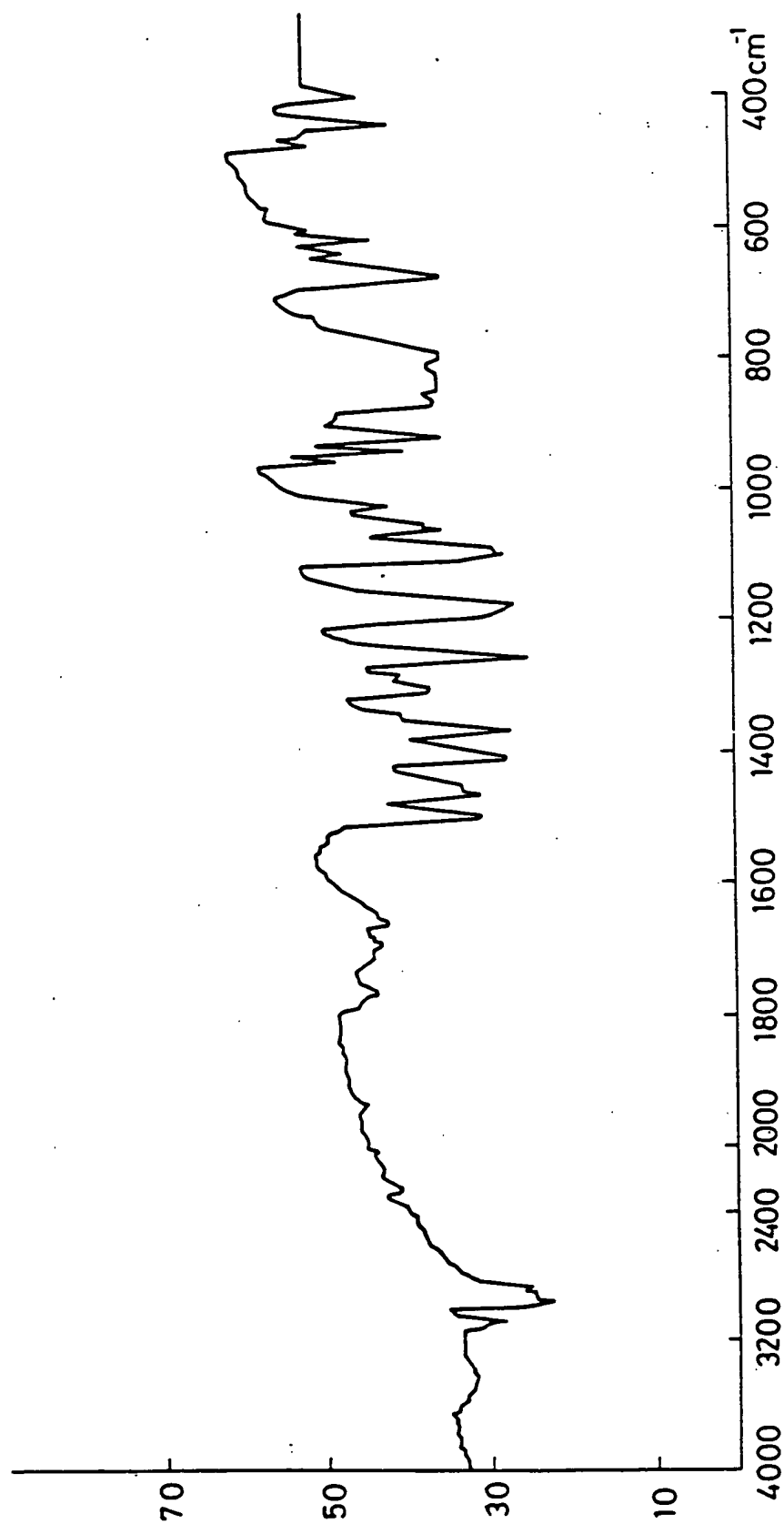


FIG. 8

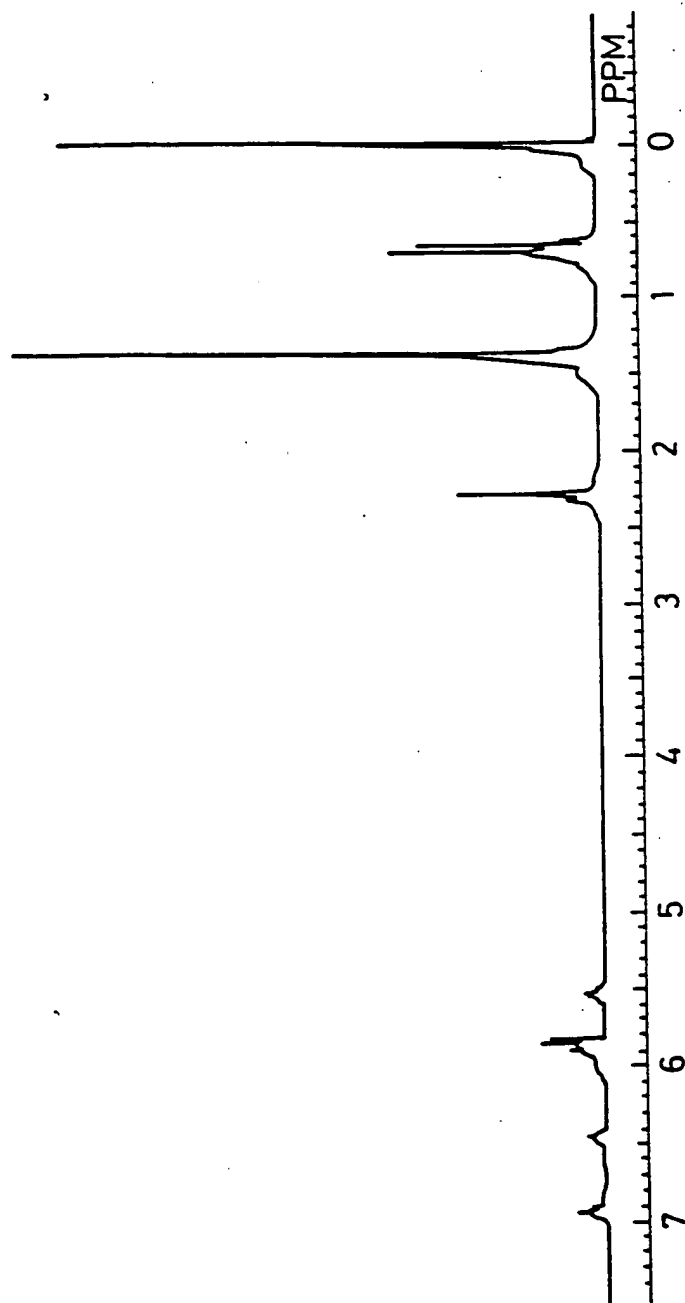


FIG. 9

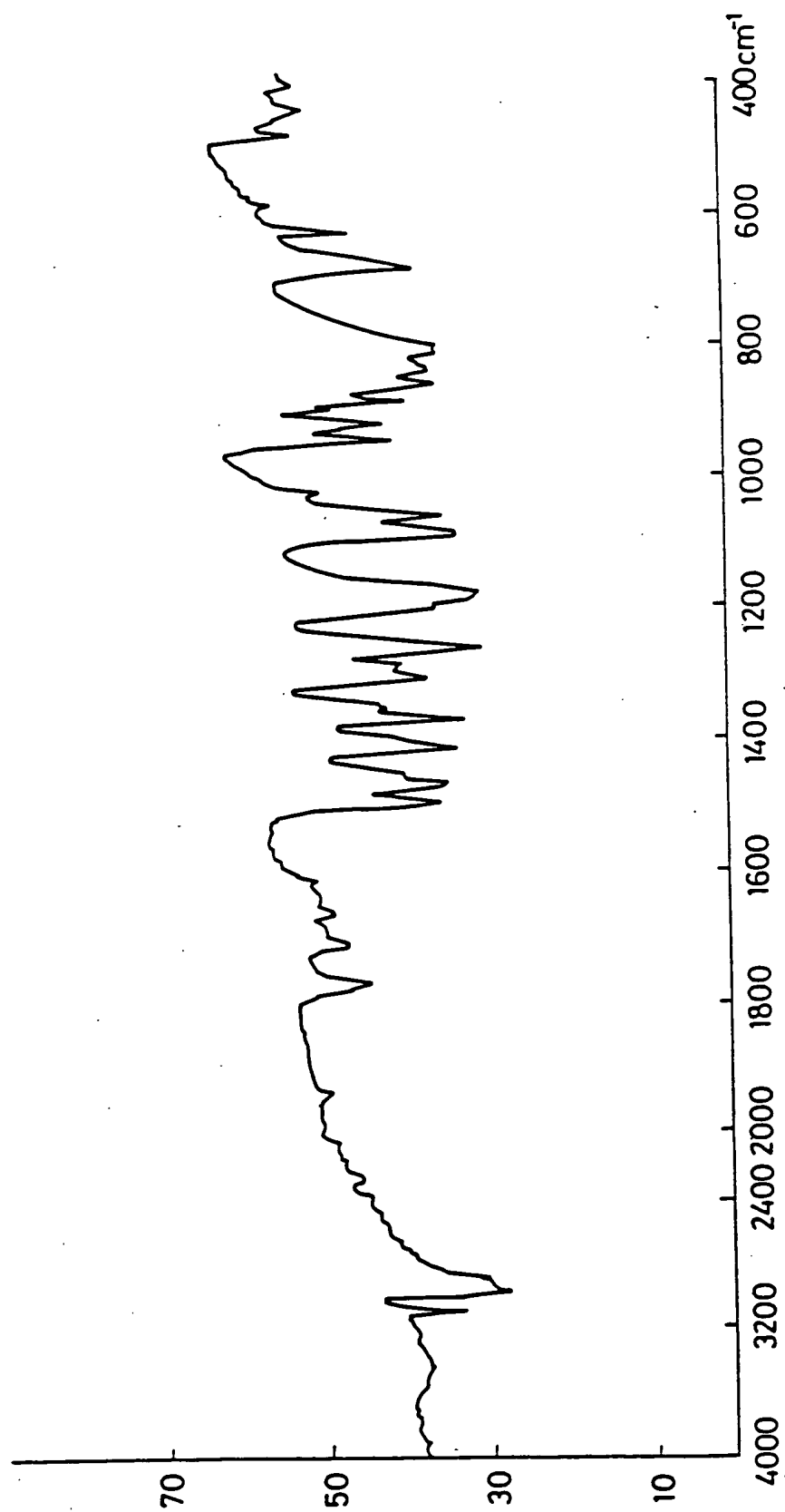


FIG.10

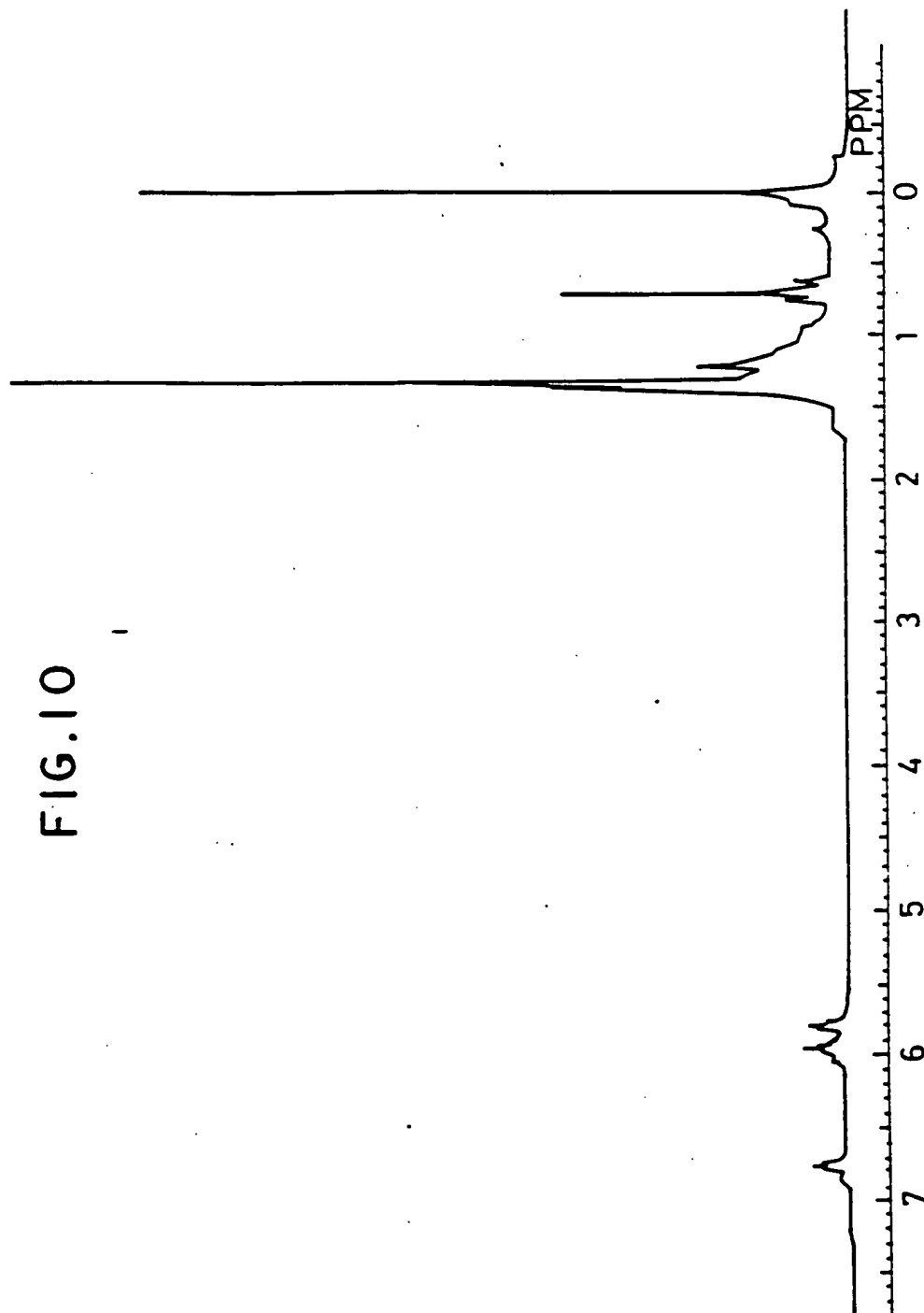


FIG. 11

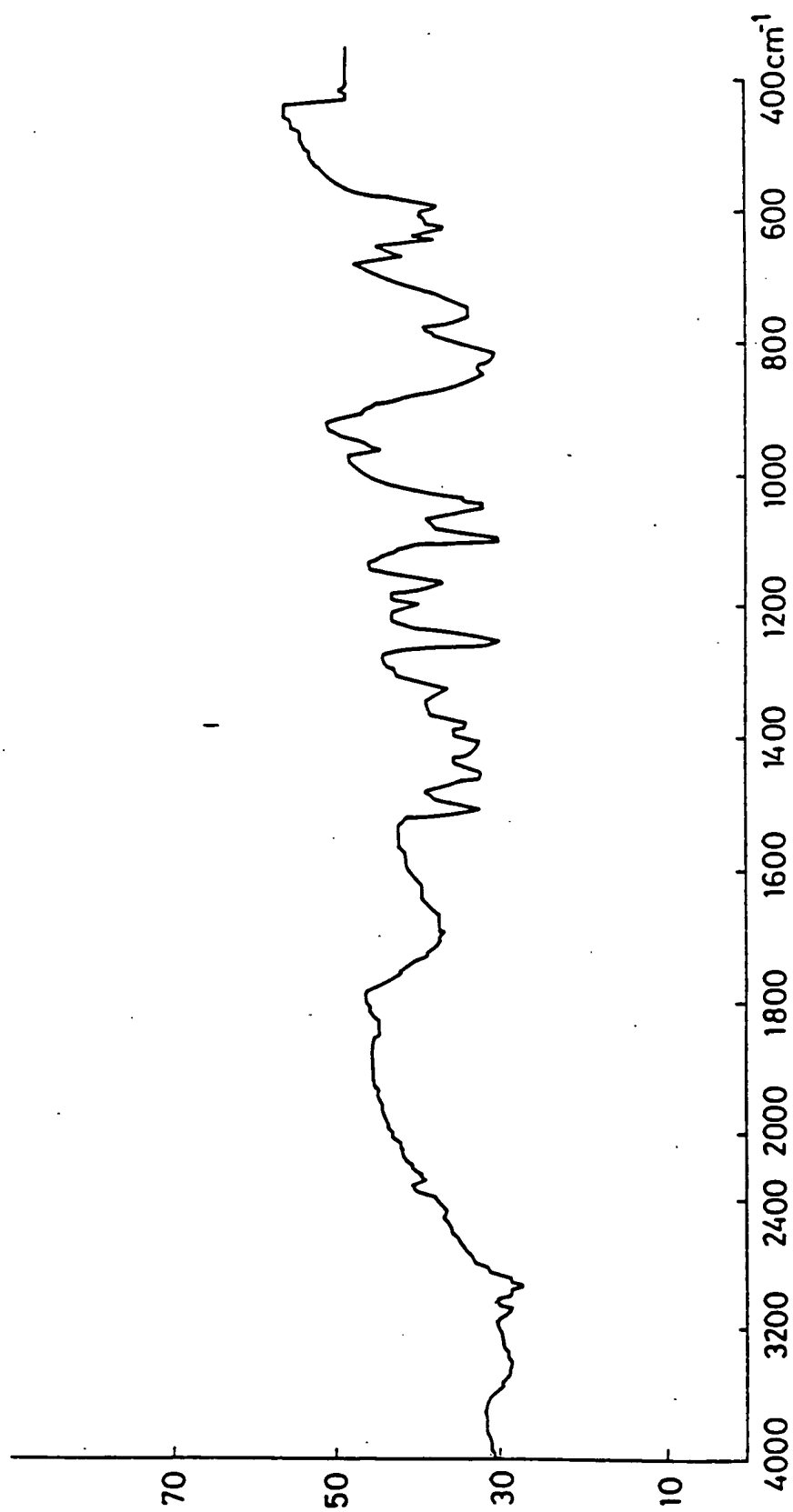


FIG. 12

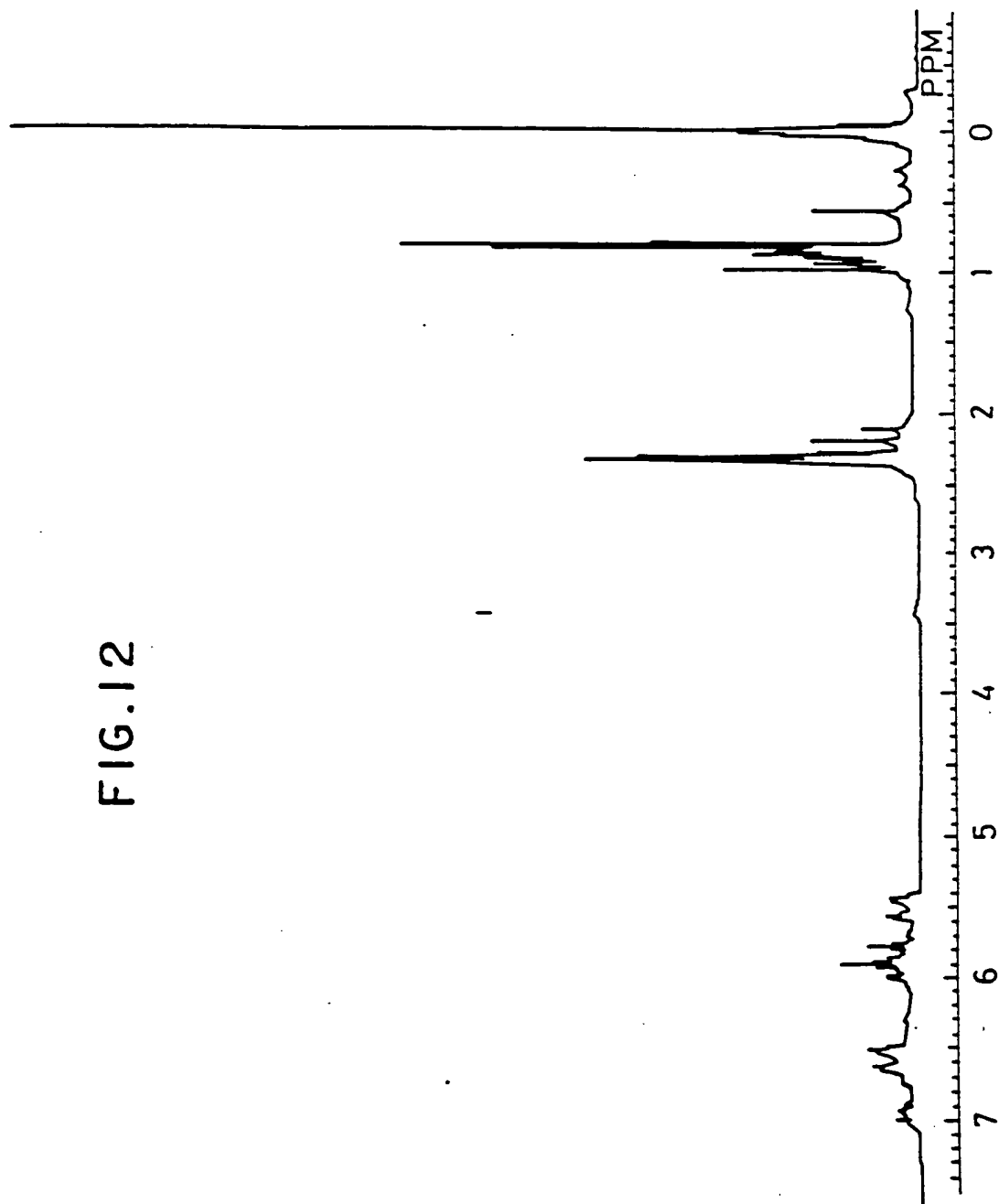


FIG. 13

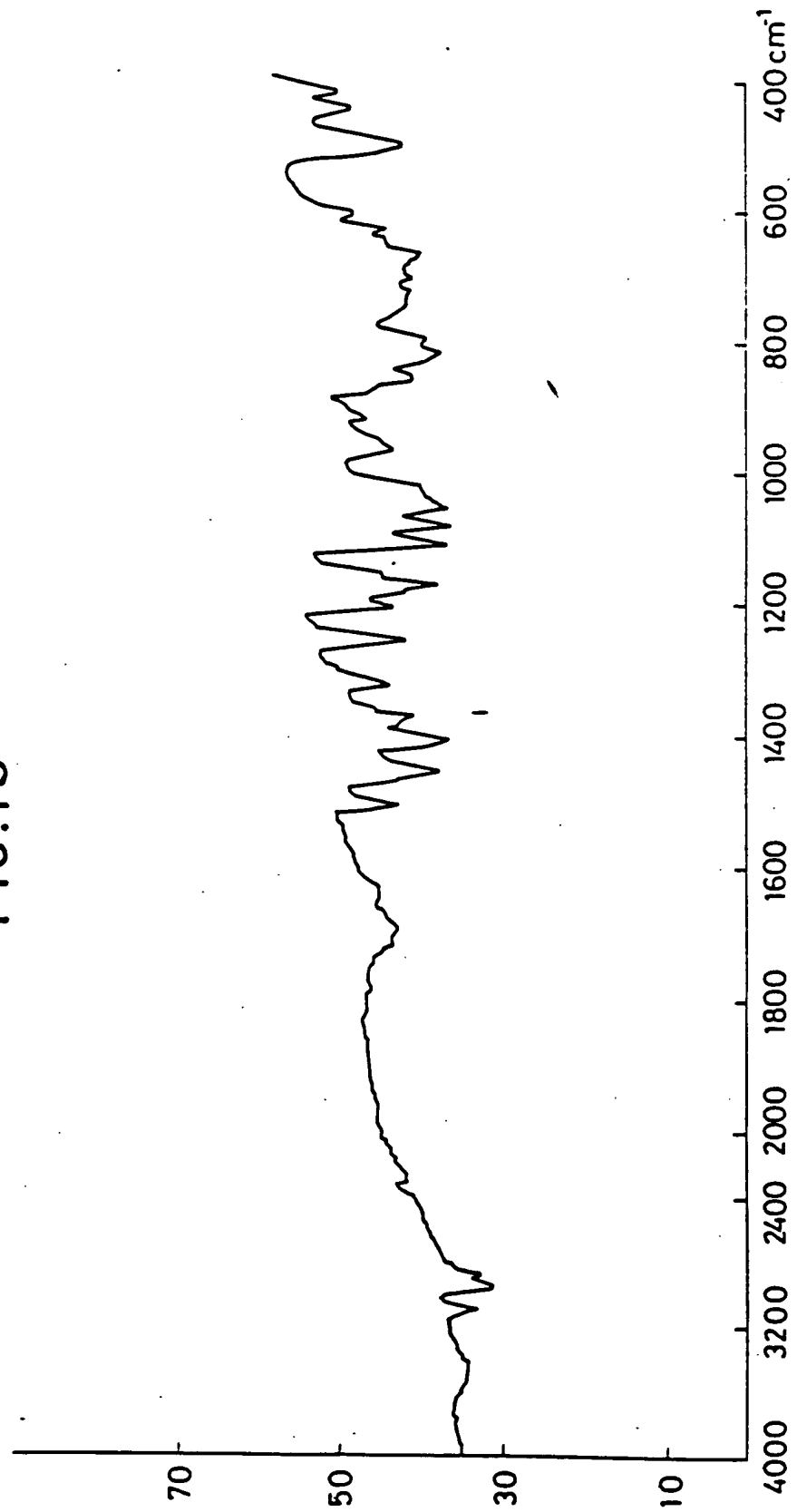


FIG. 14

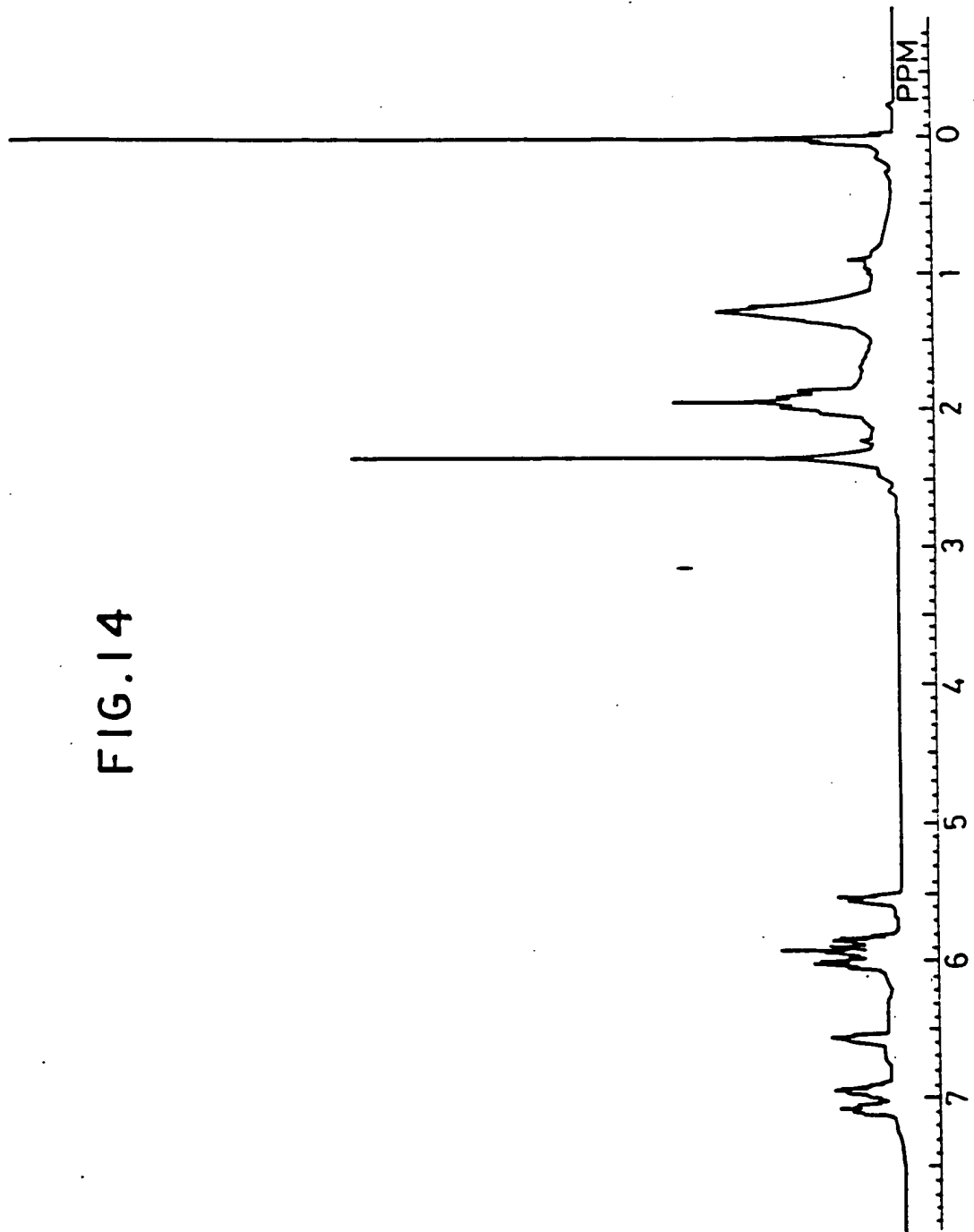


FIG. 15

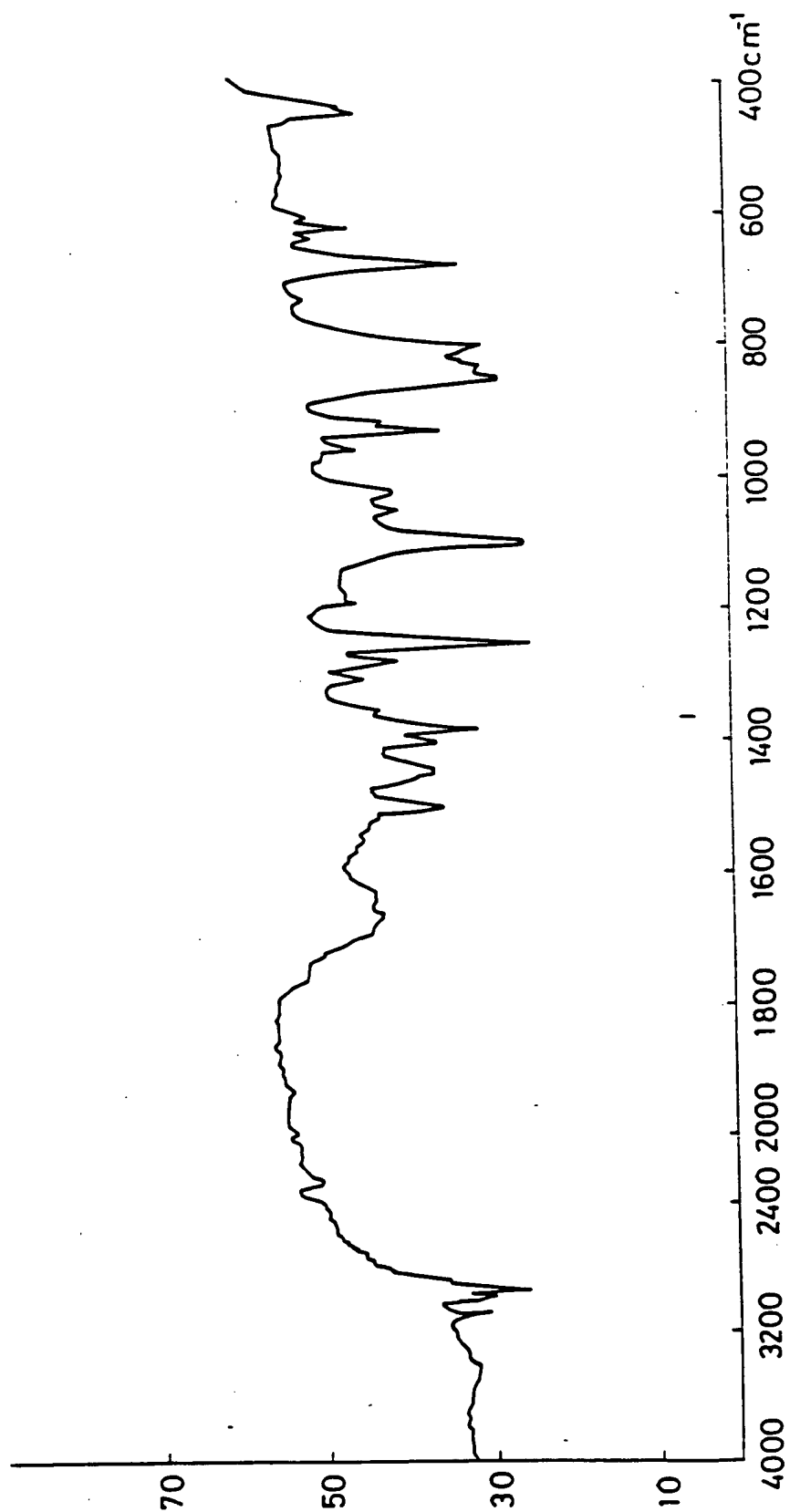


FIG.16

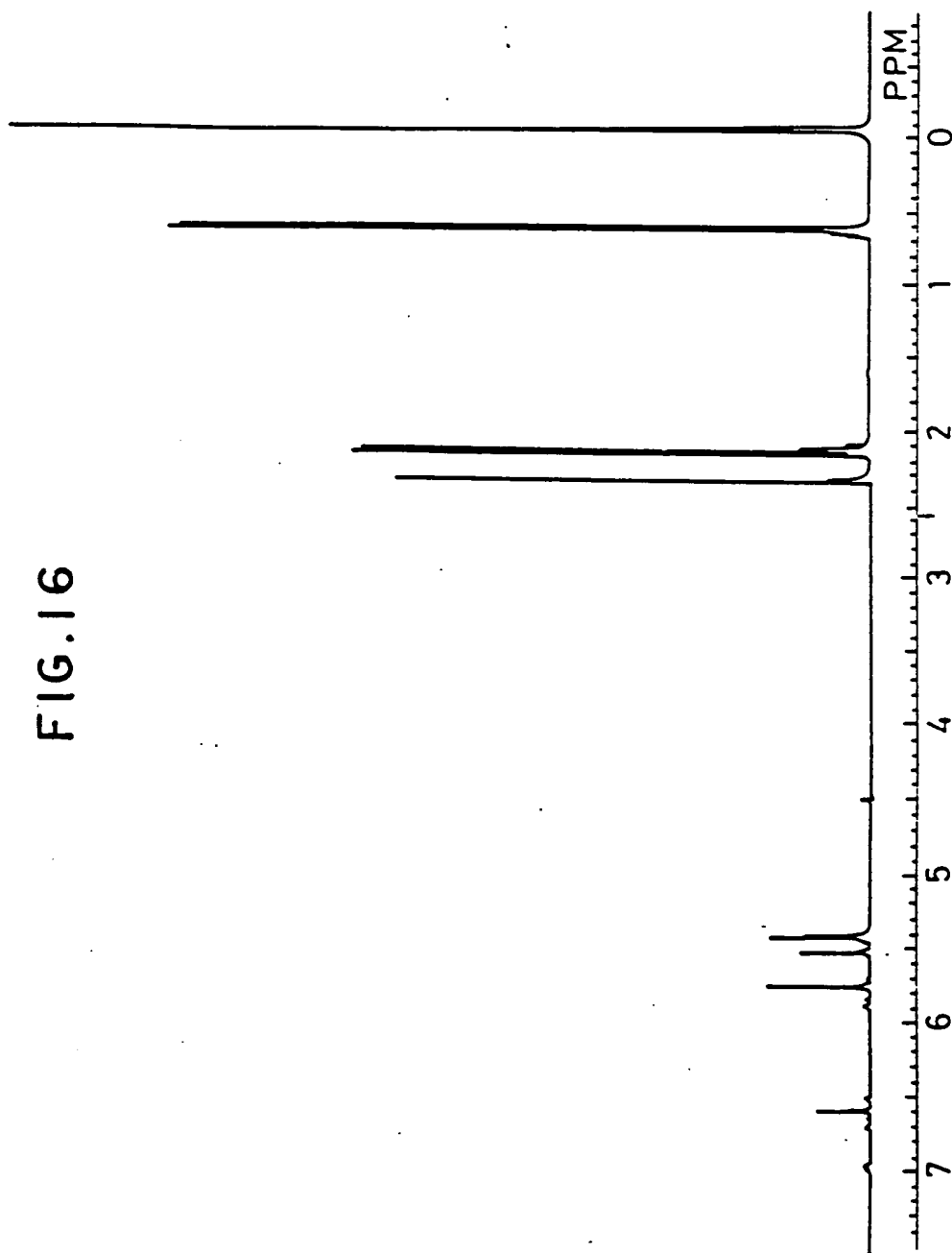


FIG.17

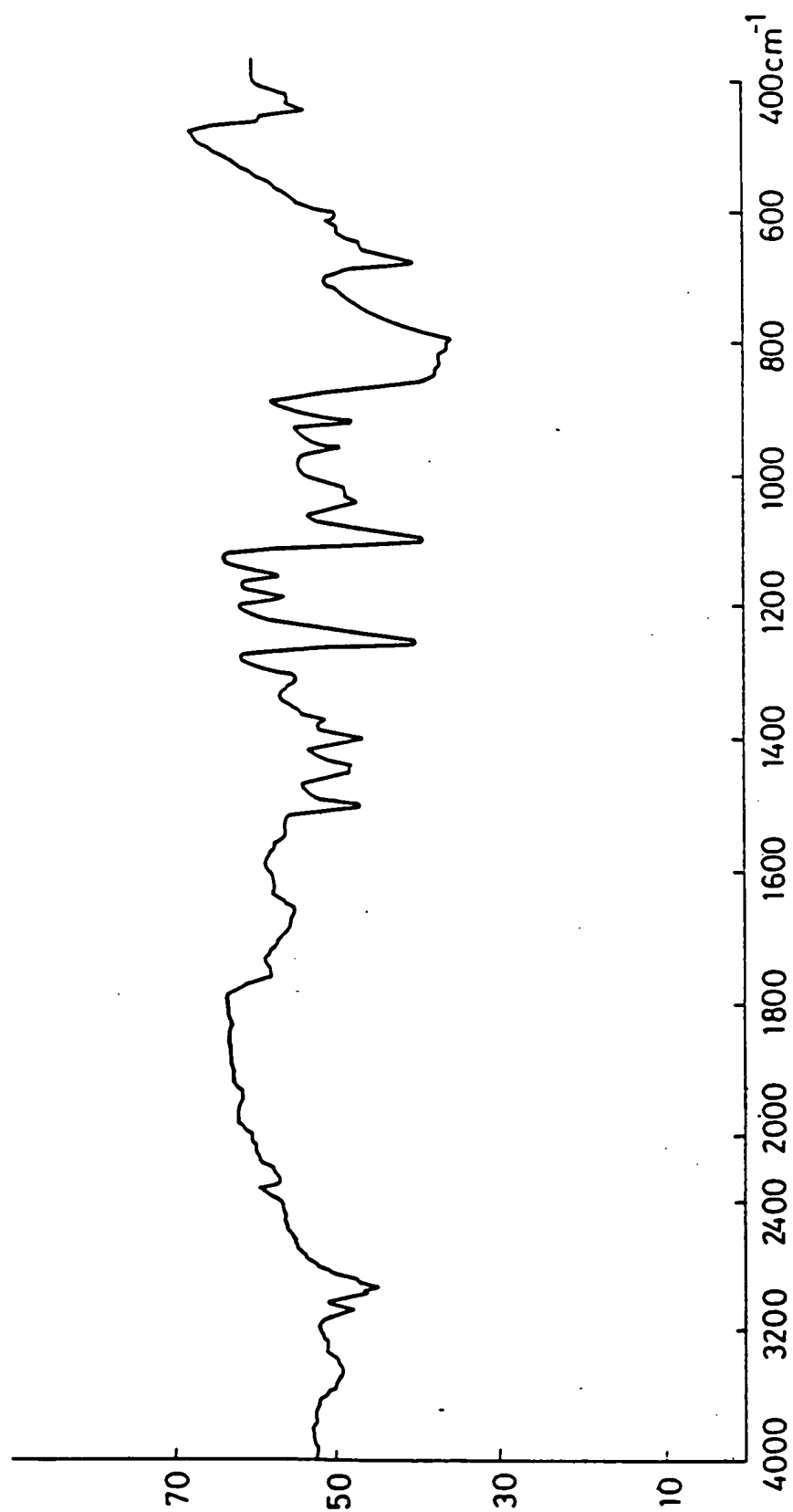


FIG. 18

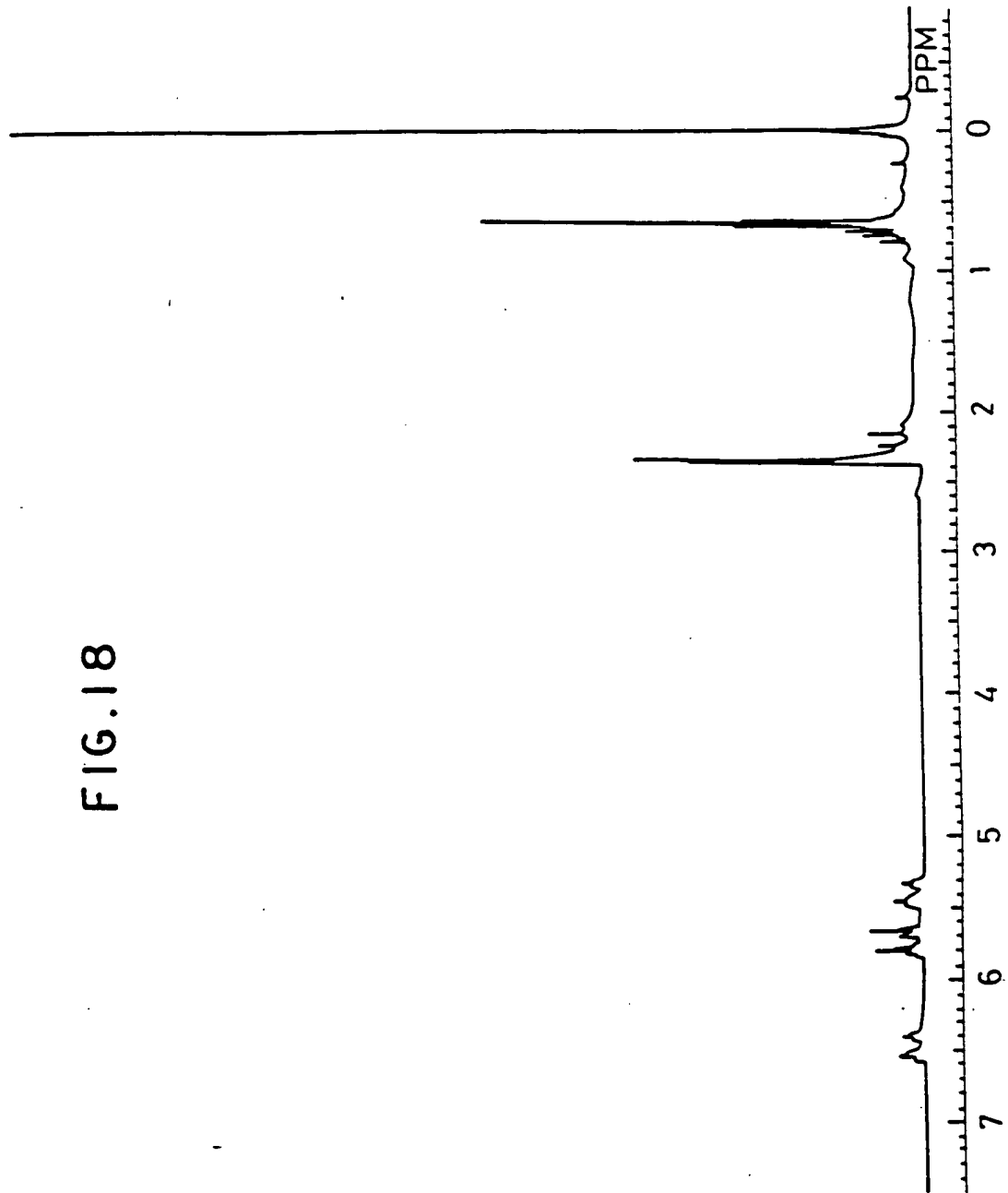


FIG. 19

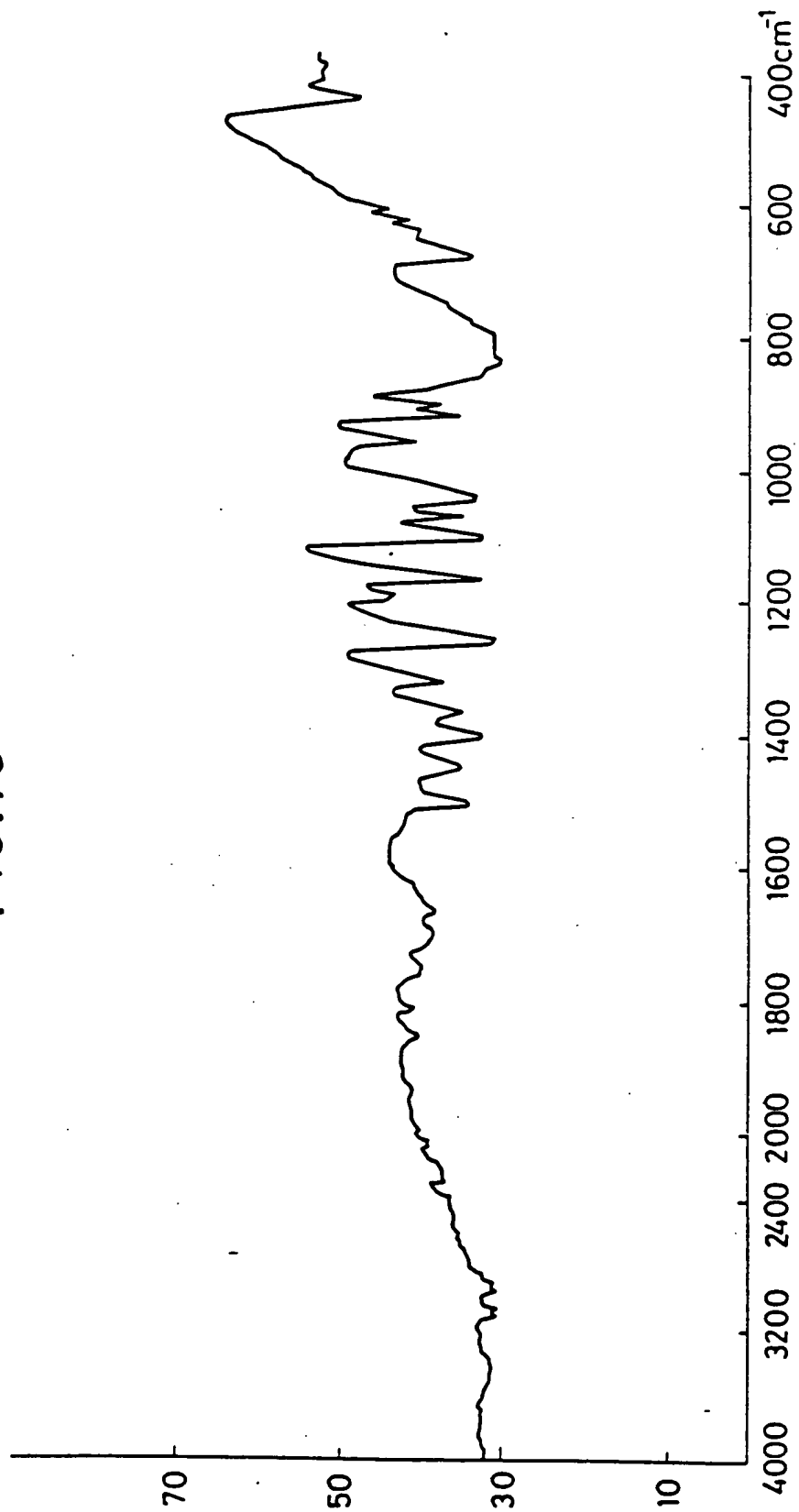


FIG. 21

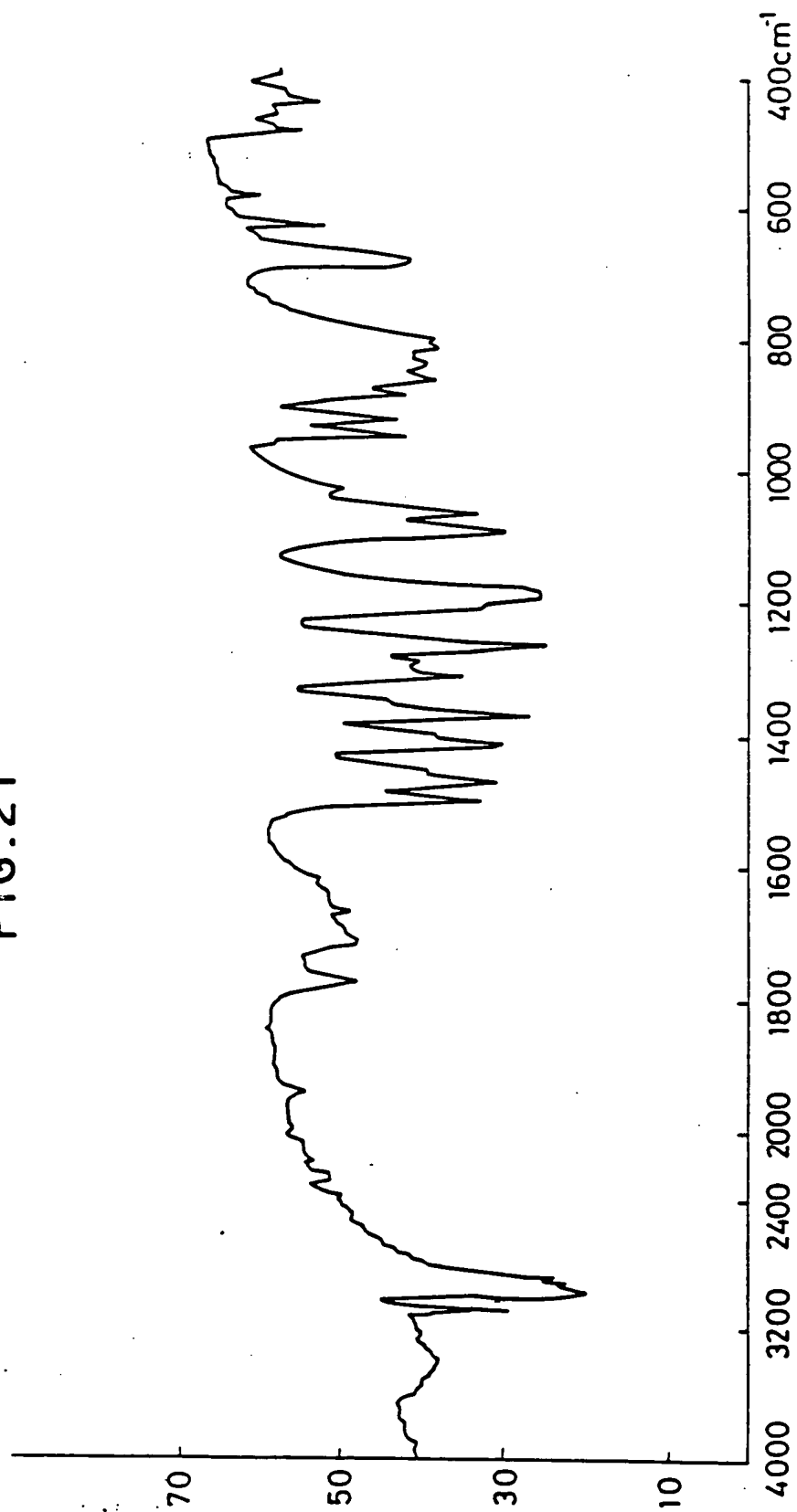


FIG. 20

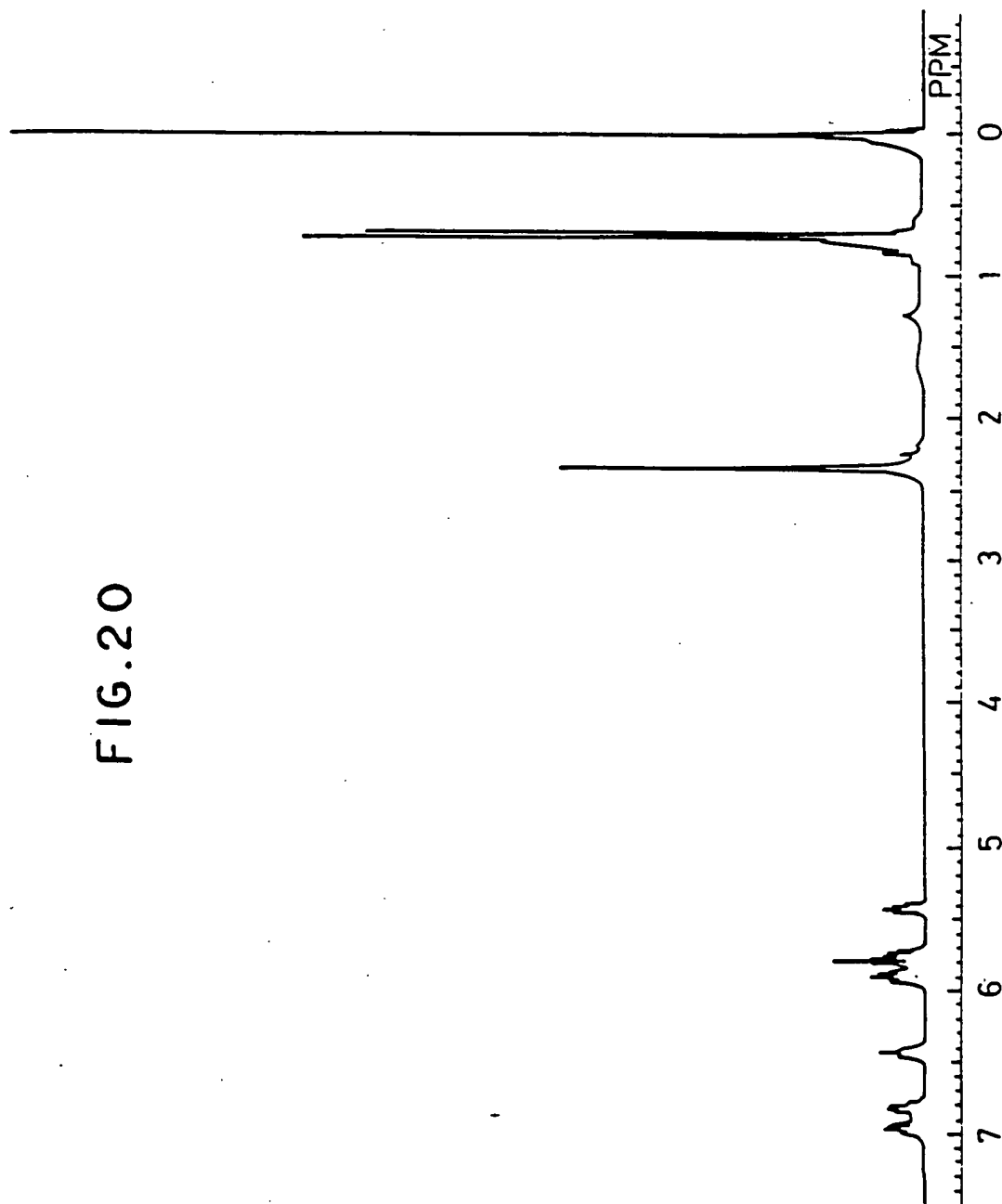


FIG. 22

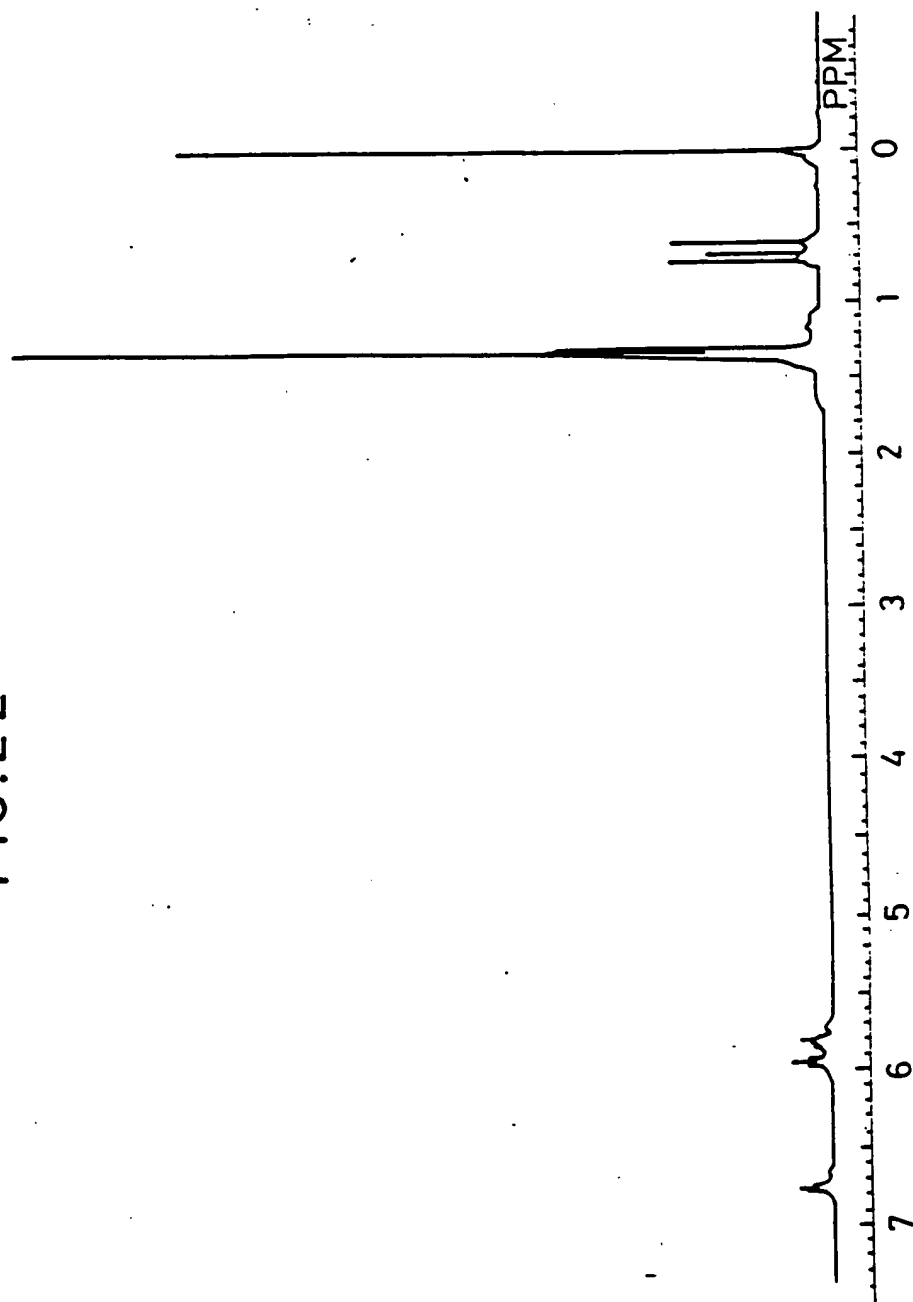
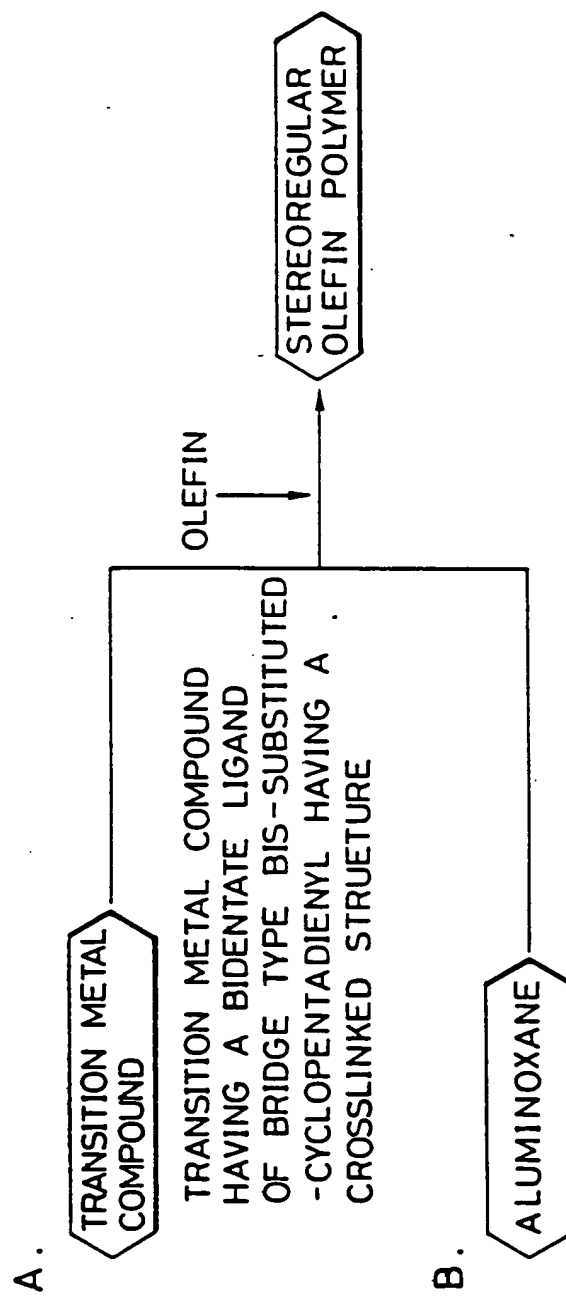


FIG. 23



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